

# *The Student's Physics*

## MODERN PHYSICS

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# MODERN PHYSICS

BY

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## PREFACE

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This book is based largely on lectures which I have given to students taking an Honours Course in Physics during recent years. It is intended to serve as an introduction to the treatises and papers on particular branches of the subject to several of which reference is made at the end of each chapter. It is impossible for any student to acquire a complete knowledge of all branches of physics, and in this book I have tried only to give a concise but intelligible account of no more than a serious student of physics ought to be familiar with when he begins to specialize on some particular branch.

The rapid changes which are now taking place in several of the subjects discussed make it difficult to write a consistent and up-to-date account of them.

Very few references to original papers have been given. These will be found in the treatises on special branches mentioned at the end of each chapter.

No attempt has been made to assign all the facts and theories discussed to their original authors. This has been done only in a few cases. The particular facts and theories selected for discussion in such a book depend largely on the knowledge and experience of the writer, and it is impossible to avoid many omissions of important results some of which another author would very likely have considered it essential to include.

My best thanks are due to Dr. John Dougall for his careful

reading of the proofs and for many valuable suggestions. Also to Sir Ernest Rutherford and Mr. Blackett of the Cavendish Laboratory for permission to reproduce the beautiful photographs of  $\alpha$ -ray tracks.

H. A. WILSON.

RICE INSTITUTE, HOUSTON, TEXAS,

*May, 1928.*

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# MODERN PHYSICS

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## CHAPTER I

### The Electron Theory

#### 1. Electrons and Positive Nuclei.

The principal electrical properties of matter are specific inductive capacity, conductivity, and magnetic permeability. When the numerical values of these quantities for the bodies present in a system are known, then the electrical phenomena to be expected in the system may be worked out. For most practical purposes it is sufficient to know that different forms of matter have these properties in greater or less degree, and it is not necessary to attempt to explain why this is so.

For scientific purposes, however, it is desired to explain the properties of matter, and to do this it is necessary to consider the nature of the ultimate particles composing material bodies. The properties, such as those just mentioned, which can be measured, are properties of matter in bulk or of bodies containing enormous numbers of atoms. They represent average values taken over volumes very large compared with the volume of one atom.

According to the electron theory material bodies contain enormous numbers of minute particles of negative electricity which are all equal. These atoms of negative electricity are called electrons. Electrically neutral bodies contain equal amounts of positive and negative electricity, and the positive electricity is also supposed to consist of minute particles which are the nuclei of the atoms.

The electrons and positive nuclei are supposed to be so small that even in the densest solids such as gold they only occupy a very minute fraction of the space. Material bodies therefore are merely space with particles of electricity here and there separated by distances very large compared with the dimensions of the particles. The space between the particles is not empty, for it is filled with the electrical and magnetic fields excited by the particles. Thus if we could select a point at random

inside a material body and examine the state of the space at that point we should almost always find nothing except an electric and a magnetic field.

The point would be in a vacuum and the specific inductive capacity, conductivity, and permeability at the point would have the usual values for a perfect vacuum. If the point happened to be inside an electron or a positive nucleus, then we may suppose that there would be a certain density of electric charge at the point in addition to the electric and magnetic fields.

## 2. Fundamental Electromagnetic Equations of the Electron Theory.

The electromagnetic equations for the interior of matter according to the electron theory are the equations for space containing nothing but electricity. The only quantities which are required in these equations are density of electricity or charge per unit volume, electric field strength, and magnetic field strength. There is no magnetism on this theory, and magnetic fields are supposed to be produced only by the motion of electricity. Following H. A. Lorentz, to whom the development of the electron theory is largely due, we shall use Heaviside's rational units and suppose that the force between two charges  $e_1$  and  $e_2$ , at rest at a distance  $r$  apart, is equal to  $e_1 e_2 / 4\pi r^2$ . This amounts to taking for the unit charge a quantity of electricity equal to the usual electrostatic unit divided by  $\sqrt{4\pi}$ . The field strength will as usual be the force on a unit charge, so that the field strength due to a charge  $e$  at rest is  $e/4\pi r^2$ . The number of lines of force crossing unit area, drawn perpendicular to the field, will, also as usual, be taken equal to the strength of the field. The number of lines of force ending on a negative charge  $e$  or starting from a positive charge  $e$  is therefore  $4\pi r^2 \times \frac{e}{4\pi r^2}$ , which is equal to  $e$ . The number of lines of force starting in unit volume is therefore equal to the density of charge, which is expressed by the equation

$$\text{div } \mathbf{F} = \rho,$$

where  $\mathbf{F}$  denotes the field strength and  $\rho$  the density of charge. The force between two magnetic poles will be taken to be  $m_1 m_2 / 4\pi r^2$ , so that the field strength due to a pole  $m$  is  $m/4\pi r^2$ . Since we suppose that there is no magnetism all lines of magnetic force must be closed curves, so that the number of lines which end in unit volume is zero, and  $\text{div } \mathbf{H} = 0$ , where  $\mathbf{H}$  denotes the magnetic field strength.

The current density, that is the current per unit area drawn perpendicular to the direction of the current, at a point between the electrons and positive nuclei will be simply the displacement current

\* The vector notation and nomenclature used here is explained at p. 334.

and so will be equal to  $\partial \mathbf{F} / \partial t$ . At a point inside an electron or a nucleus where the density of charge is  $\rho$  there will be also, if the charge is moving, a convection current density equal to  $\rho \mathbf{V}$ , where  $\mathbf{V}$  is the velocity of the electricity. The total current density at any point is therefore the resultant or vector sum of  $\partial \mathbf{F} / \partial t$  and  $\rho \mathbf{V}$ , or  $\frac{\partial \mathbf{F}}{\partial t} + \rho \mathbf{V}$ .  $\mathbf{F}$  and  $\rho$  being expressed in the Heaviside electrostatic units, the unit of current density will of course be one Heaviside electrostatic unit of charge per square centimetre per second. The magnetic field  $\mathbf{H}$  is excited by the current according to the well-known relation that the work required to take a unit pole once round a current is equal to  $4\pi$  times the current, the field and current being expressed in ordinary electromagnetic units. In vector notation,  $\text{curl } \mathbf{H} = 4\pi \mathbf{i}$ , where  $\mathbf{i}$  is the current density. If  $\mathbf{i}$  is expressed in Heaviside electrostatic units and  $\mathbf{H}$  in Heaviside electromagnetic units this becomes  $\text{curl } \mathbf{H} = \mathbf{i}/c$ , where  $c$  is the velocity of light, so that we have

$$\text{curl } \mathbf{H} = \frac{1}{c} \left( \frac{\partial \mathbf{F}}{\partial t} + \rho \mathbf{V} \right).$$

In the same way the well-known relation between the induced electromotive force round a closed curve and the rate of variation of the magnetic field through the curve may be written

$$\text{curl } \mathbf{F} = - \frac{1}{c} \frac{\partial \mathbf{H}}{\partial t}.$$

The four equations

$$\begin{aligned} \text{div } \mathbf{H} &= 0, \quad \text{div } \mathbf{F} = \rho, \\ \text{curl } \mathbf{H} &= \frac{1}{c} \left( \frac{\partial \mathbf{F}}{\partial t} + \rho \mathbf{V} \right), \quad \text{curl } \mathbf{F} = - \frac{1}{c} \frac{\partial \mathbf{H}}{\partial t}, \end{aligned}$$

are the fundamental electromagnetic equations of the electron theory. If  $\rho = 0$ , they reduce to the electromagnetic equations for space devoid of ponderable matter, or ether, as it is sometimes called.

These equations theoretically enable  $\mathbf{H}$  and  $\mathbf{F}$  to be calculated when  $\rho$  and  $\mathbf{V}$  are given throughout space as functions of the time  $t$ .

The velocity  $\mathbf{V}$  of the electricity is supposed to be measured relatively to the material system on which the observer is. In all ordinary cases  $\mathbf{V}$  will be the velocity relative to the earth or to the laboratory in which the observer is working. The field strengths  $\mathbf{F}$  and  $\mathbf{H}$  also are the fields as measured by an observer relative to the material system on which he is working, which is usually the earth. It is found that the motion of the earth has no observable influence on electromagnetic phenomena so that it is customary in electrical experiments to regard the laboratory as at rest.

### 3. Remarks on the Fundamental Equations and on the Fields.

The fundamental electromagnetic equations indicate that each charge produces a field as though the other charges were not present. Thus if all the charges are at rest and there is no magnetic field anywhere, then the electric field is given by  $\text{div } \mathbf{F} = \rho$ . This makes the field due to a charge  $e$  equal to  $e/4\pi r^2$ , and the potential  $e/4\pi r$ . The resultant field at any point is then the resultant of the fields due to every charge, each charge giving a field  $e/4\pi r^2$ . The total potential at any point is simply the sum of the potentials due to all the charges present. This means that the presence of material bodies in no way modifies the fields due to charges. Any charge excites the same field inside a material body as in empty space. This is not surprising when we remember that material bodies are almost entirely empty space, since the electrons and nuclei occupy only a minute fraction of the space even in the densest substances. It is possible that if a body could be obtained with such a high density that the electrons and nuclei in it filled an appreciable fraction of its volume, the fields produced in it would differ appreciably from those produced by the same charges in empty space.

The fundamental equations of the electron theory lead to a fairly satisfactory explanation of the electrical properties of material bodies containing very large numbers of electrons and nuclei. They fail, however, to explain the properties of single atoms. It is possible that they represent results which are true on the average over large numbers of electrons and time intervals not too short, but which are not true for very small numbers of electrons and very short time intervals. For example, the equation  $\text{div } \mathbf{F} = \rho$  may really only be true when  $\rho$  is the average density of charge over a volume containing a very large number of electrons and nuclei. However, in the electron theory we assume that this equation is true when  $\rho$  is the charge in an indefinitely small volume divided by that volume. The volume considered may be small compared with the volume of a nucleus even. We assume that the results of large-scale experiments on bodies containing enormous numbers of electrons and nuclei are true for microscopic phenomena even inside electrons.

The fundamental equations give the electric and magnetic fields excited in the surrounding space by electric charges. It is important to remember that electric and magnetic fields are not directly observed in any experiments. Only phenomena in material bodies are observed. The existence of fields in the space surrounding charges is assumed because phenomena can be conveniently described or explained by means of this assumption. In the electron theory we assume the existence of these fields but we do not attempt to explain how they are produced or what they consist of. We may if we like regard them as merely auxiliary mathematical quantities introduced into the theory for convenience in attempting to describe phenomena. Most physicists, however, believe that these fields really exist. It has been suggested that electric and magnetic fields are modifications of the ether, a medium filling all space. This hypothesis is not of much use; it is sufficient to suppose that the charges excite the fields in the surrounding space. If all space is filled with ether, then ether and space are the same thing, and we may as well regard electric and magnetic fields as modifications of space or as merely existing in space. The distinction between ether and space will never amount to anything until some method of removing the ether from a portion of space is discovered, and we have no reason to hope that anything of the sort will ever be possible.

The electric and magnetic field strengths  $\mathbf{F}$  and  $\mathbf{H}$  are vectors, and therefore the electromagnetic equations necessarily represent relations which are geometrically possible in vector fields. Thus if  $\mathbf{A}$  denotes a vector the components of which are continuous throughout a vector field, and  $\text{div } \mathbf{A} = 0$  everywhere, then we can always find a vector  $\mathbf{B}$  such that  $\text{curl } \mathbf{B} = \mathbf{A}$ . Hence, for example,

it is always possible to find an electric field  $\mathbf{F}$  such that  $\text{curl } \mathbf{F} = -\frac{1}{c}\dot{\mathbf{H}}$ , since  $\text{div } \mathbf{H} = 0$ , and therefore  $\text{div } \dot{\mathbf{H}} = 0$ . The fact that a varying magnetic field excites an electric field is nevertheless an experimental result which is not even suggested by the geometrical properties of vectors.

#### 4. Calculation of the Magnetic Field Strength. The Vector Potential.

In order to see how to calculate the field strengths  $\mathbf{F}$  and  $\mathbf{H}$  at any point, when  $\rho$  and  $\mathbf{V}$  are supposed given as functions of the time  $t$  throughout the surrounding space, we first eliminate  $\mathbf{F}$  from the fundamental equations. To do this we use the well-known vector equation \*

$$\text{curl curl } \mathbf{A} = \text{grad div } \mathbf{A} - \Delta \mathbf{A},$$

which is true for any vector  $\mathbf{A}$ . Hence, since  $\text{div } \mathbf{H} = 0$ , we have

$$\text{curl curl } \mathbf{H} = -\Delta \mathbf{H} = \frac{1}{c} \text{curl } (\dot{\mathbf{F}} + \rho \mathbf{V}).$$

But  $\text{curl } \mathbf{F} = -\frac{1}{c}\dot{\mathbf{H}}$ , so that  $\text{curl } \dot{\mathbf{F}} = -\frac{1}{c}\ddot{\mathbf{H}}$ , and therefore

$$\Delta \mathbf{H} - \frac{1}{c^2}\ddot{\mathbf{H}} = -\frac{1}{c} \text{curl } \rho \mathbf{V}.$$

If  $V$  is any quantity and  $\Delta V = \omega$ , then we have as the solution of this differential equation

$$V_P = -\frac{1}{4\pi} \int \frac{\omega dS}{r},$$

where  $V_P$  is the value of  $V$  at a point  $P$ ,  $r$  is the distance from an element of volume  $dS$  to  $P$ , and  $\omega$  the value of  $\omega$  in the element  $dS$ . The integral is supposed extended over all the space around  $P$  in which  $\omega$  differs from zero. If  $V$  and  $\omega$  are vectors then this equation is equivalent to three equations, one for each component, e.g.

$$V_x = -\frac{1}{4\pi} \int \frac{\omega_x dS}{r},$$

where  $V_x$  and  $\omega_x$  denote the  $x$  components of  $\mathbf{V}$  and  $\boldsymbol{\omega}$ .

If instead of  $\Delta V = \omega$ , we have

$$\Delta V - \frac{1}{c^2}\ddot{V} = \omega,$$

then the solution is

$$V_P = -\frac{1}{4\pi} \int \frac{[\omega] dS}{r},$$

where  $[\omega]$  stands for the value of  $\omega$ , not at the time  $t$  at which the value

of  $V$  at the point  $P$  is  $V_P$ , but at the earlier time  $t - r/c$ . The effect of the term  $-\ddot{V}/c^2$  is to make the field excited by  $\omega$  travel out from  $\omega$  with the velocity  $c$ .

The solution of the equation obtained above for  $\mathbf{H}$  is therefore

$$\mathbf{H} = \frac{1}{4\pi c} \int \frac{[\text{curl } \rho \mathbf{V}]}{r} dS.$$

Since the calculation of  $\mathbf{H}$  by means of this equation is rather complicated it is usual to introduce an auxiliary mathematical quantity  $\mathbf{a}$ , called the vector potential, which is given by the equation

$$\mathbf{a} = \frac{1}{4\pi c} \int \frac{[\rho \mathbf{V}]}{r} dS,$$

so that  $\mathbf{H} = \text{curl } \mathbf{a}$ . Here again the square brackets mean that the value of  $\rho \mathbf{V}$  in the element of volume  $dS$  is to be taken at the time  $t - r/c$  in order to get the value of  $\mathbf{a}$  at the time  $t$ .

### 5. Calculation of the Electric Field Strength. The Scalar Potential.

If we eliminate  $\mathbf{H}$  from the electromagnetic field equations in the same way as  $\mathbf{F}$  was eliminated we get

$$\Delta \mathbf{F} - \frac{1}{c^2} \ddot{\mathbf{F}} = \text{grad } \rho + \frac{1}{c^2} \frac{\partial}{\partial t} (\rho \mathbf{V}).$$

The solution of this is

$$\mathbf{F} = -\frac{1}{4\pi} \int \frac{[\text{grad } \rho + \frac{1}{c^2} \frac{\partial}{\partial t} (\rho \mathbf{V})]}{r} dS.$$

Instead of calculating  $\mathbf{F}$  by means of this vector equation we may introduce an auxiliary mathematical quantity  $\phi$ , called the scalar potential, which is given by

$$\phi = \frac{1}{4\pi} \int \frac{[\rho]}{r} dS,$$

and  $\mathbf{F}$  is then given by the equation

$$\mathbf{F} = -\frac{1}{c} \dot{\mathbf{a}} - \text{grad } \phi,$$

since

$$\frac{1}{4\pi} \int \left[ \frac{1}{c^2} \frac{\partial}{\partial t} (\rho \mathbf{V}) \right] \frac{1}{r} dS = \frac{1}{c} \dot{\mathbf{a}}.$$

When  $\rho$  and  $\mathbf{V}$  are known as functions of the time at all points in the

space around a point  $P$ ,  $\phi$  and  $\mathbf{a}$  can be calculated at any point near  $P$ , and then  $\mathbf{H}$  and  $\mathbf{F}$  are given by

$$\mathbf{H} = \text{curl } \mathbf{a}, \quad \mathbf{F} = -\frac{1}{c} \dot{\mathbf{a}} - \text{grad } \phi.$$

## 6. Examples of the Use of the Equations.

As an example of the use of these equations we will work out the fields due to a single element of volume in which  $\rho$  and  $\mathbf{V}$  are supposed constant. For this element of volume let  $\rho dS = e$ , so that  $\phi = e/4\pi r$  and  $\mathbf{a} = e\mathbf{V}/4\pi cr$ .

Let the element of volume considered be at the origin of co-ordinates so that  $r^2 = x^2 + y^2 + z^2$ , and let the velocity  $\mathbf{V}$  be along the  $x$  axis, so that  $V_x = V$ ,  $V_y = 0$ ,  $V_z = 0$ . Then we have

$$a_x = \frac{eV}{4\pi cr}, \quad a_y = 0, \quad a_z = 0, \quad \text{and } \dot{\mathbf{a}} = 0.$$

Hence

$$H_x = (\text{curl } \mathbf{a})_x = 0,$$

$$H_y = (\text{curl } \mathbf{a})_y = \frac{\partial}{\partial z} \left( \frac{eV}{4\pi cr} \right) = -\frac{eVz}{4\pi cr^3},$$

$$H_z = (\text{curl } \mathbf{a})_z = -\frac{\partial}{\partial y} \left( \frac{eV}{4\pi cr} \right) = \left( \frac{eVy}{4\pi cr^3} \right).$$

The resultant magnetic field  $H$  is therefore equal to  $\frac{eV}{4\pi cr^3} \sqrt{z^2 + y^2}$ , or, if we put  $\sqrt{z^2 + y^2} = r \sin \theta$ , it is equal to  $\frac{eV}{4\pi cr^2} \sin \theta$ . Its direction is perpendicular to the plane containing  $r$  and  $\mathbf{V}$ , so that the lines of magnetic force are circles with their centres on the  $x$  axis and their planes perpendicular to the  $x$  axis. The electric field  $\mathbf{F} = -\text{grad } \phi$ , since  $\dot{\mathbf{a}} = 0$ , which gives  $\mathbf{F} = \frac{e}{4\pi r^2}$ .

If we suppose that the element of volume considered contains equal amounts of positive and negative electricity so that  $\rho = 0$  in it, but that the positive electricity only has the velocity  $\mathbf{V}$ , then we get

$$\phi = 0, \quad \text{and } a_x = \frac{eV}{4\pi cr},$$

so that the element produces the resultant magnetic field  $H = \frac{eV \sin \theta}{4\pi cr^2}$ , but no electric field. The element of volume is then equivalent to an element of a conductor carrying a current, and the formula we have obtained for  $H$  gives Ampere's formula for the magnetic field due to a current element, since  $eV$  is equivalent to  $i ds$ , where  $i$  is the current and  $ds$  the length of the element.

As another example, suppose that an electron with charge  $e$  has been at rest at the origin from  $t = -\infty$  to a time  $t_1$ , and that between  $t_1$  and  $t_2$  it moves a small distance away from the origin and then back again and then remains at rest. Up to  $t_1$  the field will be  $e/4\pi r^2$  along  $r$ , but when the electron begins to move the field will change and the change will move out from the charge with velocity  $c$ . If we describe two spheres with centres at the origin and radii  $c(t - t_1)$  and  $c(t - t_2)$ , then outside the larger sphere the disturbance due to the motion will not have arrived, and inside the smaller sphere the disturbance will have passed by, so that the field will be  $\frac{e}{4\pi r^2}$  along  $r$  except in the space between the two spheres. Inside

this space there will be a magnetic field due to the motion as well as an electric field. Thus the motion of the electron produces a wave in the field which moves out from it with the velocity  $c$ .

### 7. Poynting's Theorem.

We shall as usual suppose that the energy density in the electromagnetic field is  $\frac{1}{2}(\mathbf{F}^2 + \mathbf{H}^2)$ , so that if  $\mathbf{P}$  denotes the flow of energy through unit area in unit time, then

$$-\operatorname{div} \mathbf{P} = \frac{1}{2} \frac{d}{dt} (\mathbf{F}^2 + \mathbf{H}^2) = (\mathbf{F} \cdot \dot{\mathbf{F}}) + (\mathbf{H} \cdot \dot{\mathbf{H}}).$$

In a space where  $\rho = 0$  we have

$$\dot{\mathbf{F}} = c \operatorname{curl} \mathbf{H}, \quad \text{and} \quad \dot{\mathbf{H}} = -c \operatorname{curl} \mathbf{F},$$

so that we get

$$\begin{aligned} -\operatorname{div} \mathbf{P} &= c(\mathbf{F} \cdot \operatorname{curl} \mathbf{H}) - c(\mathbf{H} \cdot \operatorname{curl} \mathbf{F}) \\ &= -c \operatorname{div} [\mathbf{F} \cdot \mathbf{H}]. \end{aligned}$$

Hence we conclude that  $\mathbf{P} = c [\mathbf{F} \cdot \mathbf{H}]$ , a result first obtained by Poynting and known as Poynting's Theorem. According to this the flow of energy is perpendicular to both the electric and magnetic fields and equal per unit area per unit time to  $cFH \sin \theta$ , where  $\theta$  is the angle between  $\mathbf{F}$  and  $\mathbf{H}$ .

### 8. Electromagnetic Momentum.

We know that there is a force on a current in a magnetic field due to the interaction of the magnetic field of the current and the external field. In space where  $\rho = 0$  the current density is  $\dot{\mathbf{F}}$ , so that there must be a force on the electromagnetic field equal to  $\frac{1}{c} [\dot{\mathbf{F}} \cdot \mathbf{H}]$  per unit volume.

Also there must be a similar force on a varying magnetic field in an electric field, for a varying magnetic field produces an electric field in exactly the same way that a current produces a magnetic field.

This force will be  $\frac{1}{c} [\mathbf{F} \cdot \dot{\mathbf{H}}]$  per unit volume.

The total force on the field is therefore  $\frac{1}{c} [\dot{\mathbf{F}} \cdot \mathbf{H}] + \frac{1}{c} [\mathbf{F} \cdot \dot{\mathbf{H}}]$ , which is equal to  $\frac{1}{c} \frac{d}{dt} [\mathbf{F} \cdot \mathbf{H}]$  or  $\dot{\mathbf{P}}/c^2$  per unit volume. Since force is equal to rate of change of momentum we conclude that the field has momentum equal to  $\mathbf{P}/c^2$  per unit volume. This momentum is called the electromagnetic momentum of the field. Its existence was first pointed out by J. J. Thomson. We do not regard this momentum as momentum of the ether but as momentum of the electromagnetic field. Thus the field has energy and momentum and can move through space so that

it has the most essential properties of matter. We regard the field as excited in space by the charges and it is not necessary to introduce the idea of an ether into the discussion.

Since it appears that the momentum is equal to  $\mathbf{P}/c^2$ , we conclude that the momentum is due to the flux of energy  $\mathbf{P}$ , so that we may say that electromagnetic energy when moving has momentum. The momentum of matter may therefore be supposed to be simply the momentum of the energy it contains. For energy can be converted from one kind into another, so that if electromagnetic energy has momentum it is difficult to see how other kinds of energy could not have the same amount of momentum.

### 9. Momentum and Energy of Matter.

Matter perhaps contains other kinds of energy besides electromagnetic energy, but we shall suppose that all the energy in matter has the same momentum as electromagnetic energy. If we make this assumption we can easily find the momentum of any material system in terms of its energy.

Let  $\bar{x}$  denote the  $x$  co-ordinate of the centroid of the energy in any system, so that

$$\bar{x} = \frac{\int E x dS}{\int E dS},$$

where  $E$  is the energy density in the element of volume  $dS$ , and the integrals are supposed taken over the whole of the system. Putting  $\mathcal{E} = \int E dS$  for the total energy we get

$$\mathcal{E} \bar{x} = \int E x dS.$$

Differentiating this with respect to the time  $t$  we get

$$\mathcal{E} \dot{\bar{x}} = \int \dot{E} x dS;$$

but  $\dot{E} = -\operatorname{div} \mathbf{P}$ , so that

$$\mathcal{E} \dot{\bar{x}} = -\int x (\operatorname{div} \mathbf{P}) dS.$$

Now 
$$x \operatorname{div} \mathbf{P} = \frac{\partial}{\partial x} (x P_x) + \frac{\partial}{\partial y} (x P_y) + \frac{\partial}{\partial z} (x P_z) - P_x,$$

and 
$$\int \left\{ \frac{\partial}{\partial x} (x P_x) + \frac{\partial}{\partial y} (x P_y) + \frac{\partial}{\partial z} (x P_z) \right\} dS$$

is, by Green's Theorem, equal to a surface integral over a surface enclosing the volume  $S$ . If all the energy is inside this closed surface then

$\mathbf{P}$  is zero all over it, so that the integral must be equal to zero. Hence

$$\mathcal{E}\dot{\bar{x}} = \int P_x dS.$$

But  $P_x = M_x c^2$ , where  $M_x$  is the  $x$  momentum density, so that if  $\mathcal{M}_x = \int M_x dS$  we get

$$\mathcal{M}_x = \frac{\mathcal{E}\dot{\bar{x}}}{c^2}.$$

Thus the momentum of any system is equal to its energy multiplied by the velocity of the centroid of its energy and divided by the square of the velocity of light.

#### 10. Relations between the Energy, Momentum, Velocity, and Mass of a Particle.

For a small particle of any kind the velocity  $v$  of the particle may be put equal to the velocity of the centroid of its energy, so that for any particle

$$\mathcal{M} = \frac{\mathcal{E}v}{c^2}.$$

If a force  $f$  acts on such a particle along the direction in which it is moving, and we suppose that there is no loss of energy by radiation or otherwise, then we have

$$\begin{aligned} f\delta t &= \delta\mathcal{M}, \\ f\delta x &= \delta\mathcal{E}. \end{aligned}$$

But  $\delta x = v\delta t$ , so that  $v\delta\mathcal{M} = \delta\mathcal{E}$ .

The equation  $\mathcal{M} = \mathcal{E}v/c^2$  gives  $\delta\mathcal{M} = \frac{\mathcal{E}\delta v + v\delta\mathcal{E}}{c^2}$ ,

so that

$$\delta\mathcal{E} = \frac{\mathcal{E}v\delta v + v^2\delta\mathcal{E}}{c^2}$$

or

$$\frac{\delta\mathcal{E}}{\mathcal{E}} = \frac{v\delta v}{c^2 - v^2}.$$

Integrating, we obtain

$$\log \mathcal{E} = -\frac{1}{2} \log(c^2 - v^2) + \log \mathcal{E}_0 + \frac{1}{2} \log c^2,$$

where  $\mathcal{E}_0$  is the value of  $\mathcal{E}$  when  $v = 0$ .

Hence

$$\mathcal{E} = \frac{\mathcal{E}_0}{\sqrt{1 - v^2/c^2}}.$$

Since the momentum is equal to  $\mathcal{E}v/c^2$  we have for the momentum of the particle

$$\mathcal{M} = \frac{\mathcal{E}_0 v}{c^2 \sqrt{1 - v^2/c^2}}.$$

Let us now consider the mass of the particle. This we define in such a way that when a mass  $m$  has speed  $v$  its momentum is  $mv$ . The value just found for  $\mathcal{M}$  therefore gives

$$m = \frac{\mathcal{E}_0}{c^2 \sqrt{1 - v^2/c^2}},$$

or, putting  $m_0 = \mathcal{E}_0/c^2$ ,

$$m = \frac{m_0}{\sqrt{1 - v^2/c^2}}.$$

The mass  $m$  is sometimes called the apparent mass of the particle. It appears that  $m$  increases with  $v$  and becomes infinite when  $v = c$ . It is therefore impossible for the velocity of a material system to be as great as the velocity of light  $c$ . The kinetic energy of the particle is

$$\mathcal{E} - \mathcal{E}_0 = \mathcal{E}_0 \left\{ \frac{1}{\sqrt{1 - v^2/c^2}} - 1 \right\}.$$

When  $v/c$  is very small this gives

$$\mathcal{E} - \mathcal{E}_0 = \mathcal{E}_0 \left\{ 1 + \frac{v^2}{2c^2} - 1 \right\} = \frac{\mathcal{E}_0 v^2}{2c^2},$$

or, since  $m_0 = \mathcal{E}_0/c^2$ , we have  $\mathcal{E} - \mathcal{E}_0 = \frac{1}{2}m_0 v^2$ .

## 11. Relation of the Mass of an Electron to its Charge.

If the particle considered is an electron,  $m$  will be the mass of the electromagnetic field which it excites and which moves along with it, together with any additional mass which it may have. If the electron is merely an electric charge it may have no additional mass, but if it has some internal energy besides its electrical energy it will have some additional mass corresponding to this additional energy. In any case its mass should vary with its velocity in accordance with the expression found above for  $m$ , since this should hold for a particle of any kind. The experiments of Kaufmann, Bucherer, and others on the variation of the mass of electrons with their velocity have shown that the mass does vary approximately in accordance with the above formula. These experiments confirm the idea that momentum is due to flux of energy, but they give no information as to the constitution of electrons.

If we assume that an electron consists merely of a charge  $e$  uniformly distributed over the surface of a sphere of radius  $a$ , then its electrical energy when at rest is  $\frac{e^2}{8\pi a}$ , so that its mass  $m_0$  should be  $\frac{e^2}{8\pi a c^2}$ . However, such a sphere of electricity would tend to fly apart owing to the repulsion

between the different parts of its charge. To counteract this repulsion we may suppose that there is a tension inside it equal to the repulsion or to  $\frac{1}{2}\left(\frac{e}{4\pi a^2}\right)^2$ . This tension may indicate the presence of an amount of energy equal to the volume of the sphere multiplied by the tension or to  $\frac{1}{2}\left(\frac{e}{4\pi a^2}\right)^2 4\pi a^3 = \frac{e^2}{24\pi a}$ .

According to this the total energy will be

$$\frac{e^2}{24\pi a} + \frac{e^2}{8\pi a} = \frac{e^2}{6\pi a},$$

so that the mass

$$m_0 = \frac{e^2}{6\pi ac^2}.$$

From the known values of  $e/m_0$  and  $e$  we can calculate  $a$ , which comes out about  $10^{-13}$  cm. We have reasons for believing that electrons are very small, but there is of course no justification for the idea that an electron is a sphere of electricity. Electricity as we know it consists of electrons, and a part of an electron if it could be examined would very likely be found to have properties quite different from those of electricity. We might as well suppose that a part of an atom of, say, helium would have the properties of a helium atom as that part of an electron or atom of electricity would have the properties of an electron. The density of electricity  $\rho$  which appears in the electromagnetic equations is supposed in the electron theory to be the charge in an element of volume divided by the volume of the element, and it is assumed that the element of volume considered may be inside an electron and small compared with the electron. Such assumptions are not really justified; all experiments relate to volumes enormously large compared with electrons, and we are really only justified in supposing that  $\rho$  in the electromagnetic equations stands for such a quantity as

$\frac{ne}{V}$ , where  $n$  is the number of electrons, each having a charge  $e$ , contained in a small volume  $V$ .

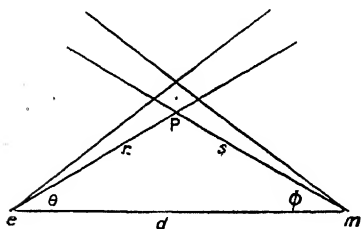


Fig. 1

## 12. Force on a Charge moving in a Magnetic Field.

The force on a charge when moving in a magnetic field can be obtained by considering the electromagnetic momentum due to a charge and a magnetic pole. Consider a charge  $e$  at a distance  $d$  from a pole of strength  $m$  (fig. 1).

Consider a point  $P$  at a distance  $r$  from  $e$  and  $s$  from  $m$ . The electric field at  $P$  due to  $e$  is  $\frac{e}{4\pi r^2}$ , and the magnetic field is  $\frac{m}{4\pi s^2}$ . The electromagnetic momentum at  $P$  per unit volume is therefore  $\frac{em \sin(\theta + \phi)}{16\pi^2 r^2 s^2 c}$ , where  $\theta$  is the angle  $Pem$  and  $\phi$  the angle  $Pme$ . As element of volume let us take a ring element described by the element of area bounded by the lines  $\theta, \theta + d\theta, \phi, \phi + d\phi$ , when the plane containing  $\theta$  and  $\phi$  rotates once round the line  $em$  as axis. The element of area is equal to  $\frac{rs d\theta d\phi}{\sin(\theta + \phi)}$ , so that the ring element of volume is equal to  $2\pi r \sin\theta \frac{rs d\theta d\phi}{\sin(\theta + \phi)}$ . The angular momentum in the ring element about the line  $em$  as axis is therefore

$$\frac{s \sin\phi em \sin(\theta + \phi) 2\pi r \sin\theta rs d\theta d\phi}{16\pi^2 r^2 s^2 c \sin(\theta + \phi)} = \frac{em \sin\theta \sin\phi d\theta d\phi}{8\pi c}.$$

The total angular momentum will be got by integrating with respect to  $\theta$  from  $\theta = 0$  to  $\theta = \pi - \phi$ , and then with respect to  $\phi$  from  $\phi = 0$  to  $\phi = \pi$ . This gives for the total angular momentum  $\frac{em}{4\pi c}$ , a result first obtained by J. J. Thomson, to whom this interesting method of calculating the force on a charge moving in a magnetic field is due. Now suppose the charge is moving with velocity  $v$  in a direction making an angle  $\psi$  with the magnetic field. In a time  $dt$  the line joining  $e$  and  $m$  will turn through an angle  $\frac{v \sin\psi dt}{d}$ , so that the angular momentum is changed in direction, and this requires the addition of angular momentum about an axis in the plane containing  $e, m$ , and  $v$ , equal to

$$\frac{emv \sin\psi dt}{4\pi cd}.$$

If a force  $f$  acts on the charge in a direction perpendicular to the plane containing  $e, m$ , and  $v$ , and an equal and opposite force on the pole, then we may equate the couple to the rate of change of the angular momentum, so getting

$$fd = \frac{emv \sin\psi}{4\pi cd}.$$

Hence

$$f = \frac{emv \sin\psi}{4\pi cd^2}.$$

If we put  $H = \frac{m}{4\pi d^2}$ , this becomes

$$f = \frac{Hev \sin\psi}{c},$$

so that for the force on a unit charge moving with velocity  $\mathbf{V}$  we get  $\frac{1}{c}[\mathbf{V} \cdot \mathbf{H}]$ . The total force on unit charge in the electromagnetic field is therefore the resultant or vector sum of the force due to the electric field and that due to the magnetic field, or

$$\mathbf{F} + \frac{1}{c}[\mathbf{V} \cdot \mathbf{H}].$$

This equation is of equal importance with the other fundamental electromagnetic equations.

### 13. Force on a Moving Electron.

When an electron having the charge  $e$  is moving with velocity  $\mathbf{V}$  in an electromagnetic field, the force on it due to the external fields  $\mathbf{F}$  and  $\mathbf{H}$  will be

$$e\left(\mathbf{F} + \frac{1}{c}[\mathbf{V} \cdot \mathbf{H}]\right),$$

provided we suppose that  $\mathbf{F}$  and  $\mathbf{H}$  do not vary appreciably in the small space occupied by the electron. In addition to this force on the electron there will be the force on it due to its own field. If the mass of the electron is zero then the total force on it must be zero, but if it contains some internal energy and so has some mass, then the total force will be equal to the rate of change of its momentum. We do not know how much internal energy there is in an electron, so we cannot tell whether its mass is zero or not.

As we have seen, the momentum of an electron or any other kind of particle is given by the expression  $\mathcal{E}\mathbf{v}/c^2$ , where  $\mathcal{E}$  is its energy. If we take the energy of the electron to be not only its internal energy but also the energy of its electromagnetic field which moves along with it, then we may consider it as a particle having momentum given by the above expression, and we may regard the forces which its own field exerts on it as internal forces which do not modify the motion of its centre of mass. The force on it may then be taken to be  $e\left(\mathbf{F} + \frac{1}{c}[\mathbf{V} \cdot \mathbf{H}]\right)$ , where  $\mathbf{F}$  and  $\mathbf{H}$  are the external electric and magnetic fields. When the velocity of the electron changes it may emit radiation, that is, a field which does not move along with it. Any such radiation field must be regarded as belonging to the external field and not to the field of the electron which moves along with it, but provided the acceleration of the electron is small the effect of the radiation emitted will be negligible. The electron may therefore be regarded as a particle of mass  $m = \frac{m_0}{\sqrt{1 - v^2/c^2}}$ , where  $m_0 = \mathcal{E}_0/c^2$  and  $\mathcal{E}_0$  is the energy of

the field of the electron which moves along with it together with its internal energy, both reckoned as when the electron is at rest.

It is important to note that the radius of an electron is not known. It is quite possible that the radius may be, say,  $10^{-10}$  cm., in which case nearly all the mass would be mass of the internal energy, and the mass of the field of the electron would be negligible. In this case the complications which are sometimes introduced by supposing the internal energy to be zero would be quite unnecessary. All the experimental results on electrons agree with the view that the electron has a charge

$e$  and mass  $m = \frac{m_0}{\sqrt{1 - v^2/c^2}}$  and speculations as to the nature of this mass beyond saying that it must be the mass of energy equal to  $mc^2$  are not justified at present.

#### 14. Radiation from an Accelerated Electron.

The radiation from an electron when its velocity varies may be calculated by means of the solution of the electromagnetic equations discussed above, but the calculation is complicated and the desired result may be obtained much more easily by a method due in principle to J. J. Thomson.

Consider an electron moving along a straight line  $AO$  with constant velocity  $v$ . At and near to  $O$  suppose the velocity changes in a short time  $\delta t$  to a constant velocity  $v'$  along  $OB$ . We suppose  $v/c$  and  $v'/c$  to be very small (fig. 2). While the electron is moving along  $AO$  its field moves with it. At  $O$  where  $v$  changes it will begin to excite the field corresponding to the velocity  $v'$  along  $OB$ , and at a time interval  $t$  later this new field will fill a sphere of radius

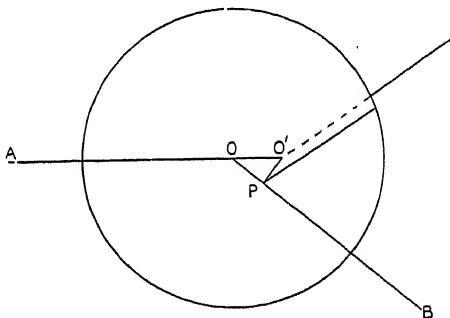


Fig 2

$ct$ . Outside this sphere the field will still be that due to the electron moving with velocity  $v$  along  $AO$ . The two fields will be separated by a layer of thickness  $c\delta t$  containing the field excited by the electron during the short interval  $\delta t$  in which its velocity changed from  $v$  to  $v'$ . This layer moves out with the velocity  $c$  and it contains the wave produced by the change from  $v$  to  $v'$ .

The lines of force in the field outside the sphere of radius  $ct$  will radiate from a point  $O'$  on  $AO$  produced such that  $OO' = vt$ , and the lines of force inside the sphere radiate from a point  $P$  on  $OB$  such that  $OP = v't$ . If we consider a line of force starting from  $P$  and making an angle  $\theta$  with  $O'P$ , it will be displaced relatively to the parallel line outside the sphere by a distance  $O'P \sin\theta$ , and we may suppose these two lines are joined into a single line by a part lying in the layer of thickness  $c\delta t$ . This requires a field component in the layer, in the plane containing  $O'P$  and parallel to the surface of the sphere, equal to

$\frac{e}{4\pi r^2} \frac{O'P \sin\theta}{c\delta t}$ , where  $r = ct$ , because the radial component in the layer is equal to  $e/4\pi r^2$ . When  $t$  is zero the line of force considered is all outside the sphere of

radius  $ct$ , and as  $t$  increases the relative displacement  $OP \sin \theta$  increases proportionally to  $t$ , so that it is clear that a line inside the sphere corresponds to a parallel line outside. Let  $OP = v''t$ , so that the tangential field in the layer is  $\frac{ev'' \sin \theta}{4\pi rc^2 \delta t}$ .

The energy in the wave per unit volume is therefore

$$\left( \frac{ev'' \sin \theta}{4\pi rc^2 \delta t} \right)^2,$$

since there must be a magnetic field of equal strength to the electric field in the wave and the energy in unit volume is  $\frac{1}{2}(E^2 + H^2)$ . The energy in the whole wave is therefore

$$\int_0^\pi 2\pi r \sin \theta r d\theta \left( \frac{ev'' \sin \theta}{4\pi rc^2 \delta t} \right)^2 c \delta t = \frac{1}{6\pi} \frac{e^2 v''^2}{c^3 \delta t}.$$

This is the energy radiated in the time  $\delta t$  while the velocity changed from  $v$  to  $v'$ , so that the rate of radiation during  $\delta t$  is

$$\frac{1}{6\pi} \frac{e^2 v''^2}{c^3 (\delta t)^2}.$$

But  $v''$  is the vector difference between  $v'$  and  $v$ , so that  $v''/\delta t$  is the acceleration of the electron during the time  $\delta t$ , and the rate of radiation of energy is

$$\frac{1}{6\pi} \frac{e^2 f^2}{c^3},$$

where  $f$  is the acceleration of the electron.

Electromagnetic radiation is obtained in practice from electrical oscillations produced by the discharge of a condenser through a wire. In such cases, in which enormous numbers of electrons are involved the radiation obtained agrees with that calculated by electromagnetic theory. Radiation from single electrons has not been observed, and according to the Quantum Theory the electrons in atoms do not radiate when they are moving round orbits and so have an acceleration. The success of the quantum theory makes it possible that the expression just obtained for the radiation from an electron is erroneous, and in fact that the equations of the electron theory are probably only true when the density of electricity  $\rho$  is taken to be the average density over a volume containing a large number of electrons and atomic nuclei. Another possibility is that the electricity in atoms is not really concentrated into small particles at all, but that the volume of an electron is comparable with the volume of an atom, so that the electrons in atoms overlap each other.

## PROPERTIES OF MATTER IN BULK

15. The electron theory gives a fairly satisfactory explanation of the principal electrical properties of matter in bulk, such as specific inductive capacity and conductivity. The specific inductive capacity ( $K$ ) of a substance may be defined as the ratio of the capacity of a condenser in which the substance fills the space between the plates to the capacity with a vacuum between the plates. For the same charges the electric field is inversely as  $K$ . The electric field  $F$  inside a material body, as for

example the insulator between the plates of a condenser, may be considered either microscopically or macroscopically. From the microscopic standpoint it is the field at a point inside the body due to all the electrons and nuclei present and it varies rapidly from point to point, being very large at points close to electrons or nuclei. From the macroscopic standpoint these variations are ignored, and we consider the field as equal to its average value over a space containing an enormous number of electrons and nuclei. Thus if  $\bar{\mathbf{F}}$  denotes this average value then

$$\bar{F}_x = \frac{1}{S} \int F_x dS,$$

where  $\bar{F}_x$  is the  $x$  component of the macroscopic field and  $F_x$  the  $x$  component of the microscopic field,  $S$  being the small volume over which the average is taken. The difference of potential between two points is given by

$$V_2 - V_1 = \int F_s ds,$$

where  $F_s$  is the microscopic field component along an element  $ds$  of the path between the two points. It is easy to see that  $\bar{V}_2 - \bar{V}_1$  is also equal to  $\int \bar{F}_s ds$ , since  $V_2 - V_1$  is independent of the path. In experimental work the field is taken to be  $\bar{\mathbf{F}}$  and is generally estimated by measuring a potential difference and dividing this by the appropriate length. The density of charge in matter may also be regarded macroscopically and we may define the macroscopic density by

$$\bar{\rho} = \frac{1}{S} \int \rho dS.$$

When  $\bar{\rho} = 0$ , what is meant is that the small volume  $S$  contains as much positive as negative electricity, or if it contains  $n$  electrons each having a charge  $e$ , and  $n'$  nuclei each having a charge  $e'$ , then  $ne + n'e' = 0$ . The electrons and nuclei in matter may be moving about so that, strictly speaking,  $\rho$  should be averaged not only over a small volume but also over a short time interval to get  $\bar{\rho}$ . The rapid fluctuations due to the motions of the electrons and nuclei are also ignored or averaged out from the macroscopic view-point. The microscopic equations  $\text{div } \mathbf{H} = 0$ ,  $\text{div } \mathbf{F} = \rho$ ,  $\text{curl } \mathbf{H} = \frac{1}{c} (\dot{\mathbf{F}} + \rho \mathbf{V})$ , and  $\text{curl } \mathbf{F} = -\frac{1}{c} \dot{\mathbf{H}}$  of the electron theory can be easily transformed into equations between the macroscopic or average values of  $\mathbf{F}$ ,  $\mathbf{H}$ ,  $\rho$ , and  $\rho \mathbf{V}$  given by

$$\bar{\mathbf{F}} = \frac{1}{S} \int \mathbf{F} dS, \quad \bar{\mathbf{H}} = \frac{1}{S} \int \mathbf{H} dS, \quad \bar{\rho} = \frac{1}{S} \int \rho dS, \quad \text{and} \quad \overline{\rho \mathbf{V}} = \frac{1}{S} \int \rho \mathbf{V} dS.$$

For a non-magnetic insulating material medium which is at rest we get

$$\operatorname{div} \bar{\mathbf{H}} = 0, \operatorname{div} \bar{\mathbf{F}} = \bar{\rho}, \operatorname{curl} \bar{\mathbf{H}} = \frac{1}{c} (\dot{\bar{\mathbf{F}}} + \bar{\rho} \bar{\mathbf{V}}), \text{ and } \operatorname{curl} \bar{\mathbf{F}} = -\frac{1}{c} \dot{\bar{\mathbf{H}}}.$$

### 16. Polarization. Electric Displacement. Equations for Non-magnetic Insulator at Rest.

The material medium is supposed to contain electrons and positive nuclei, and in the absence of an electromagnetic field the macroscopic field due to the negative charges is supposed to be equal and opposite to that due to the positive charges. When there is a field in the medium the positive charges are displaced relative to the negative charges, so that the medium acquires an electric moment or polarization  $\bar{\mathbf{P}}$ . The polarization  $\bar{\mathbf{P}}$  is defined to be the macroscopic or average electric moment per unit volume.  $\bar{\mathbf{P}}$  is a vector and we can draw lines to represent its direction. The number of lines per unit area drawn perpendicular to the lines of polarization is taken equal to  $\bar{\mathbf{P}}$ . If an electron having a charge  $e$  in the medium is displaced a distance  $\xi$  from its normal position an electric moment  $e\xi$  will be produced, since we suppose that when all the charges are in their normal positions the average moment is zero. The total moment of a small volume  $S$  will therefore be equal to  $\Sigma e\xi$ , where the sign  $\Sigma$  indicates that the products  $e\xi$  for all the positive and negative charges in  $S$  are to be summed.  $\xi$  of course is a vector, so that  $\Sigma e\xi$  indicates the vector sum of all the  $e\xi$ 's in  $S$ .

The polarization  $\bar{\mathbf{P}}$  is then equal to  $\frac{1}{S} \Sigma e\xi$ . If we draw lines of polarization through the boundary of any small area we get a tube of polarization. Consider a short length  $ds$  of such a tube and let its cross-section be  $\alpha$ , so that its volume is  $\alpha ds$  and its electric moment  $\mathbf{P} \alpha ds$ . This moment is equal to that of a positive charge  $\bar{\mathbf{P}} \alpha$  and a negative charge  $\bar{\mathbf{P}} \alpha$  at a distance  $ds$  apart. Thus we see that if a tube of polarization starts in the medium it must start from a charge  $-\bar{\mathbf{P}} \alpha$ , and if a tube ends in the medium it must end on a charge  $+\bar{\mathbf{P}} \alpha$ . The number of lines of polarization in the tube is  $\bar{\mathbf{P}} \alpha$ , so that one line of polarization starts from a unit of negative charge. We see therefore that  $\operatorname{div} \bar{\mathbf{P}} = -\bar{\rho}_P$ , where  $\bar{\rho}_P$  denotes the average charge density due to, variation of the polarization. Let  $\bar{\rho}_E$  denote the average charge density due to electricity in the medium, not due to variations in the polarization. The charge  $\bar{\rho}_E$  would remain when the field was reduced to zero but the charge  $\bar{\rho}_P$  would disappear. The equation

$$\operatorname{div} \bar{\mathbf{F}} = \bar{\rho}$$

therefore gives, since  $\bar{\rho} = \bar{\rho}_P + \bar{\rho}_E$ ,

$$\operatorname{div} (\bar{\mathbf{F}} + \bar{\mathbf{P}}) = \bar{\rho}_E.$$

Now consider the term  $\rho \bar{\mathbf{V}}$ . We have

$$\rho \bar{\mathbf{V}}_x = \frac{1}{S} \int \rho V_x dS.$$

Here  $\rho$  is the charge density in  $dS$  and  $V_x$  the  $x$  component of the velocity of the electricity. Since the electricity is supposed to consist of small particles of which a very large number are in the volume  $S$ , we may replace  $\int \rho V_x dS$  by  $\Sigma e V_x$ , which denotes the sum of the products  $e V_x$  for all the charges in  $S$ .

But  $\Sigma e V_x = \frac{d}{dt} \Sigma e \xi_x$ , where  $\xi_x$  is the  $x$  component of the displacement of a particle from its normal position. Hence, since  $\bar{P}_x = \frac{1}{S} \Sigma e \xi_x$ , we get

$$\dot{\bar{\mathbf{P}}} = \frac{1}{S} \Sigma e \mathbf{V}.$$

The equation  $\text{curl } \bar{\mathbf{H}} = \frac{1}{c} (\dot{\bar{\mathbf{F}}} + \rho \bar{\mathbf{V}})$

therefore becomes  $\text{curl } \bar{\mathbf{H}} = \frac{1}{c} (\dot{\bar{\mathbf{F}}} + \dot{\bar{\mathbf{P}}}) = \frac{1}{c} \dot{\bar{\mathbf{D}}}$ ,

where  $\bar{\mathbf{D}} = \bar{\mathbf{F}} + \bar{\mathbf{P}}$ .  $\bar{\mathbf{D}}$  is called the electric induction or the total polarization,  $\bar{\mathbf{F}}$  being the polarization of the electric field and  $\bar{\mathbf{P}}$  the additional polarization due to the matter present.  $\bar{\mathbf{D}}$  is also sometimes called the electric displacement.

The equations for the non-magnetic insulator at rest may now be written, with the dashes omitted, in the form

$$\begin{aligned} \text{div } \mathbf{H} &= 0, \quad \text{div } \mathbf{D} = \rho, \\ \text{curl } \mathbf{H} &= \frac{1}{c} \dot{\mathbf{D}}, \quad \text{curl } \mathbf{F} = -\frac{1}{c} \dot{\mathbf{H}}. \end{aligned}$$

Here,  $\mathbf{H}$ ,  $\mathbf{F}$ ,  $\mathbf{D}$ , and  $\rho$  now stand for the macroscopic average values, which can be determined experimentally.

### 17. Specific Inductive Capacity. Refractive Index.

The specific inductive capacity  $K$  of the medium may be defined as the ratio of the total polarization  $\mathbf{D}$  to the field strength  $\mathbf{F}$ , so that  $K = \mathbf{D}/\mathbf{F}$ . Hence

$$\text{curl } \mathbf{H} = \frac{1}{c} K \dot{\mathbf{F}}, \quad \text{curl } \mathbf{F} = -\frac{1}{c} \dot{\mathbf{H}}.$$

These equations give  $\text{curl curl } \mathbf{H} = -\Delta \mathbf{H} = \text{curl } \left( \frac{1}{c} K \dot{\mathbf{F}} \right)$ ,

or  $\Delta \mathbf{H} - \frac{K}{c^2} \ddot{\mathbf{H}} = 0$ .

We see therefore that electromagnetic waves will travel in the medium with the velocity  $c/\sqrt{K}$ , so that the refractive index of the medium relative to a vacuum is equal to  $\sqrt{K}$ . Since  $\mathbf{P} = \Sigma e\boldsymbol{\xi}$ , where the  $\Sigma$  now indicates the vector sum of the products  $e\boldsymbol{\xi}$  for a unit volume, we have

$$K = \frac{\mathbf{F} + \mathbf{P}}{\mathbf{F}} = 1 + \frac{\Sigma e\boldsymbol{\xi}}{\mathbf{F}}.$$

The relation between  $\Sigma e\boldsymbol{\xi}$  and  $\mathbf{F}$  may be found if we suppose that the charges in the insulator are acted on by elastic restoring forces proportional to the displacements  $\boldsymbol{\xi}$  from the normal positions. The force on a charge  $e$  will be the restoring force  $-\alpha\boldsymbol{\xi}$ , where  $\alpha$  is a constant, the force  $\mathbf{F}e$  due to the macroscopic electric field, and the force due to any field arising from the other electrons and nuclei near. The actual field at a particle is the microscopic field there, and so is not necessarily equal to  $\mathbf{F}$ . The field at the particle may be estimated by supposing that the small space around it, in which there are no other charges, may be regarded as a small spherical cavity in the medium with the particle at its centre. For the purpose of calculating the field in such a cavity we may regard the medium as consisting of a uniform distribution of positive electricity of density  $\rho$ , and a uniform distribution of negative electricity of equal density  $-\rho$ , so that the total density is zero. When the medium is polarized we suppose the positive electricity displaced relative to the negative in the direction of the polarization through a distance  $\boldsymbol{\xi}$  given by  $\mathbf{P} = \rho\boldsymbol{\xi}$ . The field due to a polarized sphere at a point inside it is then equal to the field due to a positive sphere together with that due to an equal negative sphere, the distance between the centres of the two spheres being equal to  $\boldsymbol{\xi}$ . The field due to a sphere of density  $\rho$  at a distance  $r$  from its centre is equal to  $\frac{4}{3}\pi\rho r^3/4\pi r^2$  or  $\rho r/3$ , so that the resultant field due to the two spheres is equal to  $-\rho\boldsymbol{\xi}/3$  at any point inside the two spheres. The field due to the charges on the walls of a spherical cavity in the medium is equal and opposite to that due to the sphere removed to make the cavity, so that the total field in the cavity is equal to  $\mathbf{F} + \frac{\rho\boldsymbol{\xi}}{3}$  or  $\mathbf{F} + \frac{\mathbf{P}}{3}$ . It is sometimes stated that the field due to a sphere uniformly filled with polarized atoms all equally polarized in the same direction is equal to zero, but this statement is clearly erroneous. The demagnetizing field, for example, in a uniformly magnetized sphere is well known to be equal to  $\frac{\mathbf{I}}{3}$ , where  $\mathbf{I}$  is the intensity of magnetization.

If then we suppose that a charged particle in the medium can be regarded as being inside a small spherical cavity, then the force on it will be equal to  $e(\mathbf{F} + \mathbf{P}/3)$ , due to the macroscopic field  $\mathbf{F}$  and the polarization  $\mathbf{P}$ .

In the case of a steady field  $\mathbf{F}$  we have therefore

$$e(\mathbf{F} + \mathbf{P}/3) - \alpha\boldsymbol{\xi} = 0.$$

This, with  $K = 1 + \frac{\Sigma e\boldsymbol{\xi}}{\mathbf{F}}$  and  $\mathbf{P} = \mathbf{F}(K - 1)$ , gives

$$\frac{K - 1}{K + 2} = \frac{1}{3} \frac{\Sigma e^2}{\alpha}.$$

If the electric field is not steady, then it is necessary to take into account the inertia of the charged particles. We may suppose that the nuclei remain fixed so that only the electrons have to be considered. For a particular electron of mass  $m$  and charge  $e$  we have then (dropping the vector notation)

$$m\ddot{\boldsymbol{\xi}} = -\alpha\boldsymbol{\xi} + e(\mathbf{F} + \mathbf{P}/3).$$

If we assume for simplicity that  $\alpha$  is the same for all the electrons, we get (since  $P = \Sigma e\dot{\xi}$  so that  $\ddot{P} = \Sigma e\ddot{\xi} = ne\ddot{\xi}$ )

$$nme\ddot{\xi} = -ne\alpha\xi + ne^2(F + P/3)$$

or

$$m\ddot{P} = -\alpha P + ne^2(F + P/3),$$

where  $n$  is the number of electrons in unit volume. Taking  $F = F_0 e^{ipt}$  and  $P = P_0 e^{ipt}$ , we get

$$-mp^2 P_0 = -\alpha P_0 + ne^2(F_0 + P_0/3),$$

so that

$$K = 1 + \frac{P}{F} = 1 + \frac{ne^2}{\alpha - \frac{ne^2}{3} - mp^2}.$$

Thus the specific inductive capacity and so the refractive index  $\sqrt{K}$  vary with the frequency  $p/2\pi$ . In this way a satisfactory explanation of the phenomena of dispersion can be deduced from the electron theory. By introducing a frictional force on the electrons proportional to the velocity  $\dot{\xi}$ , absorption can also be explained. These questions are more fully discussed in works on Light.

## 18. Equations for a Ferromagnetic Substance at Rest.

So far we have supposed the medium to be a non-magnetic insulator at rest. In the case of a ferromagnetic substance, on the electron theory, it is supposed that the atoms contain electrons describing orbits in such a way that the atoms have a magnetic moment, or else that the atoms contain spinning electrons which have a magnetic moment. The magnetic moments of the atoms are assumed to be practically constant, and the magnetic properties are explained by the way in which the magnetic atoms are arranged. This question is discussed in the chapter on Theories of Magnetism. Here we shall consider the macroscopic electromagnetic equations for a ferromagnetic substance at rest. In the case of the non-magnetic insulator we supposed that the charges could be regarded as neutralizing each other in the absence of an electric field. The normal position of an electron can be regarded as its average position over a time long compared with its period of revolution in its orbit, and the displacement  $\xi$  as a displacement of this average position. This rather crude procedure will evidently not suffice in the case of a ferromagnetic substance, because the magnetic moments of the atoms are produced by the orbital motion. The macroscopic value of  $\rho\mathbf{V}$  given by

$$\overline{\rho\mathbf{V}} = \frac{1}{S} \int \rho\mathbf{V} dS$$

does not include the orbital motion of the electrons, because the volume  $S$  is supposed to include a great many atoms, so that the orbital motions will cancel each other since they are as likely to be in one direction as in the opposite direction. The magnetic properties are due to micro-

scopic convection currents although the macroscopic convection current is zero.

If an electron is describing an orbit of area  $\alpha$  with frequency  $n$  the average current round the orbit is  $ne$ , so that the average magnetic moment of the orbit is  $n\alpha e/c$ . The intensity of magnetization due to such orbits is therefore given by

$$\mathbf{I} = \Sigma n\alpha e/c,$$

where the  $\Sigma$  indicates the vector sum of the moments of all the orbits in unit volume.

A uniformly magnetized bar may be compared with a closely packed bundle of solenoids each of small cross-section. The magnetic field in a uniform solenoid wound with  $n$  turns of wire per unit length is equal to  $nC/c$ , where  $C$  is the current in Heaviside electrostatic units. The magnetic moment of the solenoid is equal to  $Cn\alpha/c$ , where  $\alpha$  is the cross-section of the solenoid. The magnetic moment per unit volume of the solenoid is therefore equal to  $Cn/c$ , which is equal to the field strength in the solenoid. Thus the average field strength in a magnetized substance due to the magnetization is equal to the intensity of magnetization. The total macroscopic field strength or the magnetic induction is therefore  $\mathbf{B} = \mathbf{H} + \mathbf{I}$ , where  $\mathbf{H}$  is the field due to external sources. The induction  $\mathbf{B}$  is simply the average magnetic field, so that  $\mathbf{B} = \bar{\mathbf{H}} = \frac{1}{S} \int \mathbf{H} dS$ . The current  $\dot{\mathbf{D}}/c$ , however, does not include the microscopic orbital currents, so that a term to represent them must be added. The macroscopic equations for a magnetic insulator at rest are therefore

$$\begin{aligned} \operatorname{div} \mathbf{B} &= 0, \quad \operatorname{div} \mathbf{D} = \rho, \\ \operatorname{curl} \mathbf{B} &= \frac{1}{c} \dot{\mathbf{D}} + \operatorname{curl} \mathbf{I}, \quad \operatorname{curl} \mathbf{F} = -\frac{1}{c} \dot{\mathbf{B}}. \end{aligned}$$

The third equation is equivalent to  $\operatorname{curl} \mathbf{H} = \dot{\mathbf{D}}/c$ , where  $\mathbf{H}$  is the part of the magnetic field inside the substance not due to the local magnetization. It may be defined in the usual way as the force on a unit pole put in a long narrow cavity in the medium, the sides of the cavity being parallel to the direction of magnetization. This division of the magnetic field into two components  $\mathbf{I}$  and  $\mathbf{H}$  is arbitrary, but it is convenient, because of the difficulty of dealing with the orbital convection currents otherwise than by saying that they produce the magnetization  $\mathbf{I}$ .

We may remark here that, according to the recent new developments of the quantum theory, the charge of an electron describing an atomic orbit is regarded as distributed all round the orbit, so that the orbit is really much more nearly equivalent to a steady current of moment

$ne\alpha/c$  than it would be if the charge  $e$  were concentrated at a point.

If we assume the permeability  $\mu = \mathbf{B}/\mathbf{H}$  to be a constant, and put  $\mathbf{D} = K\mathbf{F}$ , the equations become

$$\begin{aligned}\operatorname{div} \mathbf{H} &= 0, \quad \operatorname{div} K\mathbf{F} = \rho, \\ \operatorname{curl} \mathbf{H} &= K\dot{\mathbf{F}}/c, \quad \operatorname{curl} \mathbf{F} = -\mu\dot{\mathbf{H}}/c.\end{aligned}$$

These equations give

$$\Delta \mathbf{H} = -\operatorname{curl} \operatorname{curl} \mathbf{H} = -\operatorname{curl} (K\dot{\mathbf{F}}/c),$$

so that

$$\Delta \mathbf{H} - \frac{\mu K}{c^2} \ddot{\mathbf{H}} = 0,$$

which shows that electromagnetic waves travel in the medium with velocity  $c/\sqrt{\mu K}$ , so that the refractive index is equal to  $\sqrt{\mu K}$ .

So far we have considered the substance to be an insulator. If it is a conductor, then the conduction current must be added to  $\dot{\mathbf{D}}$  in the third equation. If the conductivity is  $\sigma$ , the conduction current, which of course is a convection current or stream of electrons, is equal to  $\sigma \mathbf{F}$ .

## 19. Equations for a Moving Medium.

If the material medium is not at rest, as we have hitherto supposed it to be, then it is necessary to add new terms to the electromagnetic equations to take account of the convection currents due to the motion. If  $\mathbf{W}$  denotes the velocity of the medium then there is a convection current  $\mathbf{W}\rho$  due to the charge per unit volume. This charge  $\rho$ , as before, is the measurable charge and does not include the charge due to the polarization. The variation of the polarization  $\mathbf{P}$  gives a current  $\dot{\mathbf{P}}/c$  as when the medium is at rest, but in addition the motion of the polarization with the medium produces a magnetic field. The value of this field may be calculated by regarding the polarized medium as consisting of a series of electrical double layers, each layer having a positive charge  $S$  per unit area and a negative charge  $-S$  separated from the positive layer by a distance  $\xi$ . The moment of a layer per unit area is then  $S\xi$ , so that  $\mathbf{P} = N S \xi$ , where  $N$  is the number of layers per unit length. When the medium is moving with velocity  $\mathbf{W}$  the double layers move with velocity  $W \sin \theta$  parallel to themselves, where  $\theta$  is the angle between  $\mathbf{P}$  and  $\mathbf{W}$ . There is then a convection current  $SW \sin \theta$  per unit length along the positive layers, and  $-SW \sin \theta$  along the negative layers. These currents give a magnetomotive force in the layers equal to  $\frac{1}{c} SW \sin \theta$  in a direction perpendicular to  $\mathbf{P}$  and to  $\mathbf{W}$ . The average strength of this force is therefore  $\frac{1}{c} N \xi SW \sin \theta$ , or  $\frac{1}{c} P W \sin \theta$ . The force is therefore equal to the vector product of  $\mathbf{P}$  and  $\mathbf{W}$  into  $\frac{1}{c}$ , or  $\frac{1}{c} [\mathbf{P} \cdot \mathbf{W}]$ .

The equation

$$\operatorname{curl} \mathbf{H} = \frac{1}{c} \dot{\mathbf{D}}$$

for a non-magnetic insulator at rest becomes therefore

$$\text{curl } (\mathbf{H} - [\mathbf{P} \cdot \mathbf{W}]) = \frac{1}{c} (\dot{\mathbf{D}} + \rho \mathbf{W} + \mathbf{F}\sigma),$$

or

$$\text{curl } \mathbf{H} = \frac{1}{c} (\dot{\mathbf{D}} + \rho \mathbf{W} + \mathbf{F}\sigma + \text{curl } [\mathbf{P} \cdot \mathbf{W}])$$

for a non-magnetic conductor moving with velocity  $\mathbf{W}$ .  $\dot{\mathbf{D}}$  is the displacement current,  $\rho \mathbf{W}$  the convection current,  $\mathbf{F}\sigma$  the conduction current, and  $\text{curl } [\mathbf{P} \cdot \mathbf{W}]$  the convection current due to the motion of the polarization. The equations

$$\text{div } \mathbf{D} = \rho, \text{ div } \mathbf{H} = 0, \text{ and } \text{curl } \mathbf{F} = -\frac{1}{c} \dot{\mathbf{H}}$$

remain unchanged.

In the case of a ferromagnetic body moving with velocity  $\mathbf{W}$  we may regard the electron orbits as equivalent to magnetic doublets, and so the magnetization  $\mathbf{I}$  may be considered equivalent to a series of magnetic double layers. The motion of these double layers produces an electromotive force equal to  $-\frac{1}{c} [\mathbf{I} \cdot \mathbf{W}]$ , just as the electrical double layers give a magnetomotive force  $\frac{1}{c} [\mathbf{P} \cdot \mathbf{W}]$ . The equations for a magnetic moving body are therefore

$$\begin{aligned} \text{div } \mathbf{B} &= 0, \text{ div } \mathbf{D} = \rho, \\ \text{curl } \mathbf{B} &= \frac{1}{c} (\dot{\mathbf{D}} + \rho \mathbf{W} + \mathbf{F}\sigma + \text{curl } [\mathbf{P} \cdot \mathbf{W}]) + \text{curl } \mathbf{I}, \\ \text{curl } \mathbf{F} &= -\frac{1}{c} (\dot{\mathbf{B}} + \text{curl } [\mathbf{I} \cdot \mathbf{W}]). \end{aligned}$$

There is no magnetic convection current analogous to  $\rho \mathbf{W}$  or magnetic conduction current analogous to  $\mathbf{F}\sigma$ . Here  $\mathbf{B}$ , usually called the magnetic induction, is really the macroscopic magnetic field strength in the medium.

For an uncharged non-magnetic insulator moving in a constant electric field we have  $\dot{\mathbf{D}} = 0$ ,  $\rho \mathbf{W} = 0$ , and  $\mathbf{F}\sigma = 0$ ,

$$\text{so that} \quad \text{curl } \mathbf{H} = \frac{1}{c} \text{curl } [\mathbf{P} \cdot \mathbf{W}],$$

or

$$\mathbf{H} = \frac{1}{c} [\mathbf{P} \cdot \mathbf{W}].$$

The motion of an insulator in an electric field should therefore produce a magnetic field. This effect was detected by Röntgen by spinning a circular disc made of an insulator between the plates of a charged condenser. He obtained a magnetic field equal to that calculated.

When a material medium is moving in a magnetic field there is a force on the charges in the medium equal to  $\frac{1}{c} [\mathbf{W} \cdot \mathbf{H}]$  per unit charge, due to the motion of the charges in the magnetic field. The polarization  $\mathbf{P}$  when the velocity and the field are constant is therefore given by  $\mathbf{P} = (K - 1) (\mathbf{F} + \frac{1}{c} [\mathbf{W} \cdot \mathbf{H}])$ , because the polarization due to a field  $\mathbf{P}$  which exerts a force  $\mathbf{F}$  on a unit charge is equal to  $\mathbf{F}(K - 1)$ . The electrical displacement  $\mathbf{D} = \mathbf{P} + \mathbf{F}$  is therefore given by

$$\begin{aligned} \mathbf{D} &= (K-1)\left(\mathbf{F} + \frac{1}{c}[\mathbf{W} \cdot \mathbf{H}]\right) + \mathbf{F} \\ &= K\mathbf{F} + \frac{K-1}{c}[\mathbf{W} \cdot \mathbf{H}]. \end{aligned}$$

If  $\mathbf{F}$  is zero, then

$$\mathbf{D} = \frac{K-1}{c}[\mathbf{W} \cdot \mathbf{H}].$$

In the absence of a magnetic field an equal displacement would be produced by a field  $\mathbf{F}'$  given by

$$K\mathbf{F}' = \frac{K-1}{c}[\mathbf{W} \cdot \mathbf{H}],$$

so that

$$\mathbf{F}' = \frac{K-1}{K} \frac{1}{c} [\mathbf{W} \cdot \mathbf{H}].$$

This may be regarded as an electromotive force induced in the insulator by the magnetic field. The induced E.M.F. in a conductor moving in a magnetic field is equal to  $\frac{1}{c}[\mathbf{W} \cdot \mathbf{H}]$ , so that the induced E.M.F. in an insulator is to that in a conductor as  $K-1$  is to  $K$ . This induced E.M.F. in an insulator moving in a magnetic field was measured by the writer in a hollow cylinder of ebonite rotating in a magnetic field, and was found equal to the value indicated by the electron theory.

In the case of a magnetic insulator the displacement  $\mathbf{D}$  with  $\mathbf{F}$  zero is

$$(K-1) \frac{[\mathbf{W} \cdot \mathbf{B}]}{c} + \frac{[\mathbf{W} \cdot \mathbf{I}]}{c}.$$

This result is obtained by putting  $\mathbf{B}$  for  $\mathbf{H}$  in the expression for  $\mathbf{D}$  in a non-magnetic insulator when  $\mathbf{F}$  is zero, and adding on the displacement corresponding to the induced E.M.F. due to the motion of the electron orbits or equivalent magnetic doublets. The field  $\mathbf{F}'$  to give an equal displacement is therefore given by

$$K\mathbf{F}' = \frac{K-1}{c}[\mathbf{W} \cdot \mathbf{B}] + \frac{[\mathbf{W} \cdot \mathbf{I}]}{c}.$$

Putting  $\mathbf{I} = \frac{\mu-1}{\mu}\mathbf{B}$ , this gives

$$\mathbf{F}' = \frac{\mu K - 1}{\mu K} \frac{[\mathbf{W} \cdot \mathbf{B}]}{c}.$$

This result was verified experimentally by M. Wilson and the writer by measuring the induced E.M.F. in a cylinder consisting of small steel balls embedded in wax, rotating in a magnetic field. This medium was used because no insulator is known for which the permeability differs appreciably from unity.

## 20. Metallic Conduction.

On the electron theory some of the electrons in metallic conductors of electricity are supposed to be free to move about inside the material of the conductor. These free electrons are set in motion by an electric field, so producing a current.

The free electrons are usually supposed to be moving about inside metals like the molecules of a gas. In the absence of an electric field

as many move in one direction as in any other direction, so that on the average the current is zero. The electrons collide with the atoms, and we may suppose that the average distance between collisions or length of free path is  $\lambda$ . We may also suppose that after a collision with an atom the velocity of an electron is as likely to be in one direction as another, so that the average velocity of a large number of electrons which have all just had a collision is zero.

If there is an electric field of strength  $F$  in the metal, the force on an electron due to it is  $Fe$ , so that if there are  $n$  free electrons in unit volume then the rate at which their momentum in the  $F$  direction increases owing to the field  $F$  is  $Fen$ . Let  $\bar{u}$  denote the average value of the component of the velocity of an electron in the  $F$  direction, so that  $n\bar{u}$  is the momentum of the electrons in unit volume in the  $F$  direction. If  $V$  denotes the average value of the resultant velocity of the electrons taken without regard to direction, the number of collisions in unit volume in unit time is  $nV/\lambda$ , so that, since on the average the electrons lose their  $F$  momentum at each collision, the rate of loss of  $F$  momentum due to collisions is  $nV\bar{u}/\lambda$ . Hence in a steady state we have  $Fen = nV\bar{u}/\lambda$ , or

$$\bar{u} = \frac{Fe\lambda}{mV}.$$

The current density  $i$  is therefore given by

$$i = ne\bar{u} = \frac{Fne^2\lambda}{mV}.$$

The conductivity of the metal  $\frac{i}{F} = \frac{ne^2\lambda}{mV}$ .

This simple theory accounts for Ohm's law, since  $i \propto F$ . If we suppose that the heat conductivity of a metal is due to the free electrons, then we can easily obtain a relation between the electrical and thermal conductivities. We may suppose that the average kinetic energy  $\frac{1}{2}m\bar{V}^2$  of the electrons is proportional to the absolute temperature  $T$  as for the molecules of a gas on the kinetic theory, so that  $\frac{1}{2}m\bar{V}^2 = \alpha T$ , where  $\alpha$  is a constant. The heat capacity of the electrons in unit volume is then  $na$ . According to the kinetic theory the thermal conductivity of the electrons, supposing they conduct heat like a gas, is approximately  $\frac{1}{3}\lambda Vna^*$ , so that the ratio of the thermal to the electrical conductivity should be approximately

$$\frac{\frac{1}{3}\lambda Vna}{ne^2\lambda/mV} = \frac{2}{3}\left(\frac{\alpha}{e}\right)^2 T,$$

since  $m\bar{V}^2 = 2\alpha T$ , and  $V$  and  $\bar{V}$  do not differ much.

\* Cf. J. K. Roberts, *Heat and Thermodynamics*, Chapter XI, Section 8.

It is found experimentally that the electrical resistance of metals is roughly speaking proportional to the absolute temperature  $T$  and that the thermal conductivity is nearly independent of  $T$ , so that the ratio of the two conductivities is nearly proportional to the absolute temperature, in agreement with the theory. Also for most metals the ratio as found experimentally is not so very different from  $\frac{2}{3} \left(\frac{\alpha}{e}\right)^2 T$ .

These results and Ohm's law may be said to support the free electron theory of metallic conduction. However, there are many other experimental results which are not in agreement with this theory.

When a current is passed along a bar and one end of the bar is kept at a higher temperature than the other end, then to keep the temperature constant additional heat is required on account of the flow of electricity. Let the current be  $C$ , so that  $C/e$  electrons cross any section of the bar in unit time. If we suppose each electron has kinetic energy  $\alpha T$  then the additional heat required in unit time will be  $-C \frac{\alpha}{e} (T_2 - T_1)$ , where  $T_2 - T_1$  is the temperature difference between

the ends of the bar. The electrons move in the opposite direction to the current  $C$ , so that if the current is in the direction of rising temperature the amount of heat required will be negative. The specific heat of the electricity may therefore be said to be  $- \alpha/e$  according to this theory. Measurements of the specific heat of electricity in different metals show that it is positive for some metals and negative for others and in all cases much smaller than  $\alpha/e$ . It is therefore probable that the free electrons in metals, if there are any such, have much less kinetic energy than  $\alpha T$ . This conclusion is confirmed by the fact that the specific heats of metals are about what we should expect if the electrons do not take up any appreciable heat energy as the temperature rises.

## 21. Hall Effect.

When a bar carrying a current is put in a transverse magnetic field it is found that a small electric field is produced along lines perpendicular to the current and to the magnetic field. This is known as the Hall Effect. The mechanical force on the bar is  $Hi$  per unit volume, where  $H$  is the magnetic field strength and  $i$  the current density. This force is produced by the force on the electrons as they move across the field. The electrons in unit volume therefore receive momentum equal to  $Hi$  in unit time in the direction of the force  $Hi$ , which of course is perpendicular to  $H$  and  $i$ . Since the electrons are prevented from moving out of the bar by the attractions of the positive nuclei, an electric field is set up perpendicular to  $H$  and  $i$  which gives to the electrons momentum equal and opposite to that due to the magnetic field. In this way the force  $Hi$  is transferred from the electrons in the bar to the bar

itself. If this transverse electric field is denoted by  $F$  then we should have  $Fne = Hi$ , where  $n$  is the number of free electrons in unit volume and  $e$  is the charge on each electron. According to this,  $F$  should be in the same direction in all metals, but it is found that in some metals  $F$  is in the direction indicated by the theory and in others it is in the opposite direction. Also in liquid metals  $F$  is too small to be detected although they conduct quite well. The theory therefore fails to account for the observations on the Hall Effect.

It is found that the electrical resistance of a wire is increased when it is put in a magnetic field. The increase is nearly proportional to the square of the field strength. No satisfactory explanation of this effect is offered by the electron theory.

## 22. Modifications made by Quantum Theory.

It seems probable that the idea that metals contain free electrons moving about like the molecules of a gas is entirely erroneous. Instead we may suppose that the electrons describe stationary orbits round the atomic nuclei in accordance with the quantum theory. The energy of the electrons associated with these orbits is independent of the temperature and no radiation is emitted by the electrons in the orbits. An electron in any orbit is supposed to tend to jump into a near-by vacant orbit if the jump results in a decrease of energy. The energy difference is supposed to be emitted in the form of radiation.

Consider a metal in which there is an electric field  $X$  along the  $x$  axis. Let  $x_1$  be the  $x$  co-ordinate of any atom  $A$  and  $x_2$  that of another atom  $B$ . The energy associated with any electron orbit in  $A$  will be greater than that associated with the corresponding orbit in  $B$  by  $Xe(x_2 - x_1)$ . Thus if the orbit in  $B$  is vacant an electron in the orbit in  $A$  will tend to jump from  $A$  to  $B$  if the energy difference is positive, and the energy difference  $Xe(x_2 - x_1)$  will be emitted as radiation if the electron does jump. The electric field will therefore set up a motion of the electrons in the direction opposite to that of the field, so that there will be a current in the direction of the field. If  $\lambda$  denotes the average value of  $x_2 - x_1$ , and  $N$  the number of jumps per unit volume per unit time, then the current density  $i$  will be given by

$$i = N\lambda e.$$

The energy of the radiation emitted per unit volume per unit time will be  $NXe\lambda = iX$ , and we may suppose that the radiation is absorbed by the metal and converted into heat energy.

If there is a magnetic field  $H$  perpendicular to  $X$  then the momentum given to an electron, during a jump, by the magnetic field will be

$$He \int_{x_1}^{x_2} \dot{x} dt = He(x_2 - x_1).$$

We may suppose that this momentum is received by the atom into which the electron jumps. The momentum received by the metal per unit volume per unit time will therefore be  $NHe\lambda$ , or  $Hi$ , so that there will be a transverse force on the metal due to the magnetic field equal to  $Hi$ . The magnetic field does not affect the energy differences  $Xe(x_2 - x_1)$ , since it does no work on the electrons, so that on this theory if we assume that the jumps depend solely on the energy differences we should expect no Hall Effect and no change of resistance due to a magnetic field. The specific heat of the electricity should be zero, since the energy associated with the orbits is independent of the temperature. The very small Hall Effects, resistance changes, and specific heats of electricity which are observed in most metals are probably secondary effects due to the heterogeneous character of the crystalline structure of the metals. In liquid metals these effects are probably entirely absent. We may suppose that the probability of an electron jumping from one orbit to the corresponding vacant orbit in another atom is proportional to  $X$ , so that we may put  $N = N_1X$ , where  $N_1$  is the value of  $N$  in a field of unit strength. Hence  $i = N_1X\lambda e$ , so that the conductivity of the metal is  $N_1\lambda e$ .

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## CHAPTER II

### Theories of Magnetism

#### 1. Magnetic Moment due to Moving Electrons.

As regards their magnetic properties substances are usually classified as diamagnetic, paramagnetic, or ferromagnetic. Diamagnetic and paramagnetic bodies have constant permeabilities differing little from unity. For diamagnetic bodies the permeability is slightly less than unity, and for paramagnetic bodies slightly greater than unity. Ferromagnetic bodies have large permeabilities which vary with the magnetic field strength.

According to the electron theory the magnetic properties of all substances are attributed to the motions of the electrons which they contain. Magnetic effects due to the heavy positive nuclei are presumably negligible. The electrons may be describing orbits inside the atoms, or may be free electrons moving about between the atoms, or they may be spinning about an axis through their centres.

The magnetic moment of a small plane circuit carrying a current  $C$  is equal to the product of the area of the circuit and the current. An electron with charge  $e$ , describing a plane orbit of area  $a$  with frequency  $\nu$ , may be regarded as equivalent to a circuit of area  $a$  and current  $\nu e$ , so its magnetic moment is equal to  $a e \nu$ .\* The direction of the magnetic moment is perpendicular to the plane of the orbit.

Suppose that a substance contains  $n$  electrons per unit volume, all describing plane orbits, and that  $a$  is the area of an orbit and  $\nu$  the frequency of revolution of the electron in it, then the components of the magnetic moment of the substance per unit volume or its component intensities of magnetization parallel to rectangular axes  $x, y, z$  are

$$\Sigma a e \nu l, \Sigma a e \nu m, \Sigma a e \nu n$$

respectively, where  $l, m, n$  are the direction cosines of the normal to the plane of an orbit. It is understood that the sign  $\Sigma$  indicates the sum of the quantities like  $a e \nu l$  for all the  $n$  electrons. The magnetic moments

\* Cf. Chap. I, section 18. (In the present chapter ordinary units are used, not Heaviside units (Chap. I, section 2), and  $e$  is in electromagnetic, not electrostatic, units.

of any spinning electrons may be supposed included in this summation by making  $\alpha$  equal to the mean area described by the electricity in the electron.

## 2. Theory of Diamagnetism.

The theory of diamagnetism is closely related to that of the Zeemann Effect. In the chapter on the Quantum Theory, in connexion with the Zeemann Effect, it is shown that the equations of motion of an electron relative to fixed axes, in the absence of a magnetic field, retain very approximately the same form in the presence of a magnetic field  $H$ , provided the axes are supposed to rotate with angular velocity  $\omega = He/2m$  about the direction of the field. Here  $e$  is the charge and  $m$  the mass of an electron.

We see from this that when a magnetic field is produced in a substance in which the electrons are describing orbits in fields of force, the orbits in the field will be the same relative to axes rotating about the direction of the field with velocity  $\omega = He/2m$  as they were relative to fixed axes before the field was applied, provided that the application of the magnetic field does not move the fields of force.

In diamagnetic substances it is supposed that the atoms have zero magnetic moments in the absence of a magnetic field, so that the field does not tend to move them. A field  $H$ , along the  $x$  axis, therefore produces an intensity of magnetization along the  $x$  axis equal to  $-\Sigma a e \omega l^2 / 2\pi$ , since the frequencies are changed by  $\pm \omega l / 2\pi$ . The negative sign is correct, because an increasing magnetic field tends to induce currents which produce a field opposite to the inducing field. Hence the electrons in orbits which give a field opposite to the field  $H$  will be accelerated, and the electrons in orbits giving a field in the same direction will be retarded.

The permeability of the substance is therefore (since  $\mu H = B = H + 4\pi I$ ) given by

$$\begin{aligned}\mu &= \frac{H - \frac{1}{m} \Sigma a e^2 H l^2}{H} \\ &= 1 - \frac{1}{m} \Sigma a e^2 l^2.\end{aligned}$$

If we suppose that the normals to the orbits are distributed so that as many point one way as any other way, then the mean value of  $l^2$  will be  $1/3$ , and we have

$$\mu = 1 - \frac{e^2}{3m} \Sigma \alpha.$$

For example, a solid body having  $10^{24}$  atoms per cubic centimetre each

containing one electron orbit of area  $10^{-16}$  sq. cm. would have a permeability given by

$$\mu = 1 - \frac{e^2}{3m} 10^{24} \times 10^{-16} = 1 - 10^{-5} \text{ about.}$$

The susceptibility ( $\kappa$ ) is defined to be  $\frac{\mu - 1}{4\pi}$ , so that

$$\kappa = - \frac{e^2}{12\pi m} \Sigma \alpha.$$

The value of  $\kappa/\rho$ , where  $\rho$  is the density of the substance, is independent of the temperature for most diamagnetic bodies, and it is independent of the field strength also. The motion of the electrons in their orbits is independent of the temperature according to Bohr's quantum theory.

### 3. Magnetic Effect of Free Electrons.

If we suppose that the substance considered contains free electrons moving about inside it like the molecules of a gas, then these electrons will also produce a small diamagnetic susceptibility. The theory of this effect was given by Schrödinger, and later by the writer in a simpler form.

Let  $u, v, w$  be the velocity components of an electron parallel to the axes  $x, y, z$ . Let there be a magnetic field  $H$  parallel to the  $z$  axis. The equations of motion of an electron are then

$$m\dot{u} = Hev, \quad m\dot{v} = -Heu, \quad m\dot{w} = 0.$$

These equations give, putting  $He/m = \omega$ ,

$$u = A \sin(\omega t + \alpha), \quad v = A \cos(\omega t + \alpha),$$

where  $A^2 = u^2 + v^2$ .

If  $p$  denotes the angular momentum of the electron about the  $z$  axis we have

$$p = m(vx - uy) = m\{xA \cos(\omega t + \alpha) - yA \sin(\omega t + \alpha)\}.$$

Let  $x_1, y_1$  be the co-ordinates of the electron at the time  $t = 0$ , so that

$$x = x_1 + \frac{A}{\omega} \cos \alpha - \frac{A}{\omega} \cos(\omega t + \alpha),$$

$$y = y_1 - \frac{A}{\omega} \sin \alpha + \frac{A}{\omega} \sin(\omega t + \alpha).$$

Hence

$$p = m\{Ax_1 \cos(\omega t + \alpha) - Ay_1 \sin(\omega t + \alpha) + \frac{A^2}{\omega} (\cos \omega t - 1)\}.$$

Now suppose that there is a large number  $n$  of electrons in unit volume all moving with the same velocity  $V$  and describing free paths like the molecules of a gas. For each electron let  $t$  be the time since its last collision, so that  $x_1, y_1$  give its position at its last collision. The total angular momentum about the  $z$  axis of the  $n$  electrons will then be  $\Sigma p$ , and it is easy to see that the

sums of the first two terms in the expression for  $p$  will give zero, so that

$$\Sigma p = m \Sigma \frac{A^2}{\omega} (\cos \omega t - 1).$$

The mean value of  $\frac{mA^2}{\omega} (\cos \omega t - 1)$  during a free path of duration  $\tau$  is

$$\frac{mA^2}{\omega \tau} \int_0^\tau (\cos \omega t - 1) d\tau = \frac{mA^2}{\omega} \left( \frac{\sin \omega \tau}{\omega \tau} - 1 \right).$$

The number of free paths with times between  $\tau$  and  $\tau + d\tau$  out of a total number  $N$  is  $N e^{-\tau/T} d\tau/T$ ,\* where  $T$  is the mean value of  $\tau$ .

The average value of  $\frac{mA^2}{\omega} \left\{ \frac{\sin \omega \tau}{\omega \tau} - 1 \right\}$  is therefore

$$\int_0^\infty \frac{1}{T^2} e^{-\tau/T} \tau \frac{mA^2}{\omega} \left( \frac{\sin \omega \tau}{\omega \tau} - 1 \right) d\tau = - \frac{mA^2 \omega T^2}{1 + \omega^2 T^2}.$$

Hence

$$\Sigma p = - \frac{m \omega T^2}{1 + \omega^2 T^2} \Sigma A^2.$$

But since the velocities are as likely to be in one direction as another, the average value of  $A^2$  must be  $\frac{2}{3} V^2$ , so that

$$\Sigma p = - \frac{2}{3} \frac{nm \omega V^2 T^2}{1 + \omega^2 T^2}.$$

Putting  $\lambda = VT$ , we get

$$\Sigma p = - \frac{2}{3} \frac{nm \omega \lambda^2}{1 + \omega^2 \lambda^2 / V^2}.$$

The motion of the electrons with this angular momentum will produce a magnetic field along the  $z$  axis in the opposite direction to the field  $H$ .

Consider a bar of the metal of circular cross-section with its axis parallel to the field  $H$ . The angular momentum and magnetic field due to the motion of the electrons in this bar will be equal to those due to a current circulating round the surface of the bar in planes perpendicular to its axis. Let this current be  $i$  per unit length along the axis and let the number of electrons carrying it be  $N$  per unit area. The angular momentum of these electrons about the axis of the bar is  $2\pi r^2 N m v$ , where  $v$  is their velocity. The angular momentum per unit volume is therefore  $2\pi r^2 N m v / \pi r^2 = 2N m v$ . Also  $i = N e v$ , so that  $2N m v = 2m i / e$ .

The magnetic moment of the current is  $\pi r^2 i$ , so that the intensity of magnetization is equal to  $i$ . The angular momentum per unit volume is therefore  $2m i / e$ , where  $I$  is the intensity of magnetization. Hence

$$I = \frac{e}{2m} \Sigma p = - \frac{1}{3} \frac{ne \omega \lambda^2}{1 + \omega^2 T^2}.$$

The susceptibility  $\chi$  is therefore, since  $\omega = He/m$ ,

$$\chi = \frac{I}{H} = - \frac{1}{3} \frac{ne^2 \lambda^2}{m(1 + \omega^2 T^2)}.$$

In most cases  $\omega T$  is small compared with unity, so that approximately

$$\chi = - \frac{1}{3} \frac{ne^2 \lambda^2}{m}.$$

\* The symbol  $e$  is used in this chapter for the base of natural or Napierian logarithms, generally denoted by  $c$ .

This result is similar to the result previously obtained for electron orbits of area  $\alpha$ , since  $n\lambda^2$  is obtained instead of  $\Sigma\alpha/4\pi$ .

The diamagnetic susceptibility due to the free electrons in metals may be of the same order of magnitude as that due to the electrons describing orbits, but there is no way of estimating the relative values of the two.

A series of interesting papers on the properties of diamagnetic bodies has been published by Oxley.\* He finds that there is a decrease of susceptibility of about 5 per cent on passing from the liquid to the solid state. To explain this he supposes that there is a molecular field in crystalline solids due to the regular arrangement of the atoms which modifies the electron orbits, so that they are not the same as in the liquid state. This molecular field is estimated to be equivalent to a magnetic field of several million gauss, and it serves to explain the tensile strength of solids and the change of density on solidification. For details the original papers may be consulted. Cp. section 8.

#### 4. Paramagnetism. The Magnetron.

Paramagnetism is explained by supposing that the atoms have permanent magnetic moments. The magnetic moment of an atom is the vector sum of the moments of its electron orbits and spinning electrons. When this is zero in the absence of an external field the substance is diamagnetic, and when not zero it is paramagnetic. The effect of the magnetic field on the electron orbits which produces diamagnetism must also occur in paramagnetic substances, but it is usually too small to be appreciable compared with the paramagnetic effect of the permanent moments of the atoms.

It has been supposed that the atoms, electron orbits, spinning electrons, or other magnetic units supposed to be present in magnetic bodies always have moments which are exact multiples of an atomic unit of magnetic moment which is called a magneton. Several magnetons have been suggested, and many experimental results have been published which seem to support the view that magnetons exist. However, very accurate results are necessary to prove that a quantity is always a multiple of a definite atomic unit. This is especially the case when rather large multiples of the unit may occur. The idea of the magneton seems to be of doubtful value as far as can be judged at present.

#### 5. Langevin's Theory of a Paramagnetic Gas.

A theory of a paramagnetic gas was worked out by Langevin in 1905, and this theory has been the basis of most of the subsequent theoretical work on magnetism. Langevin supposed each molecule of the gas to have a fixed magnetic moment  $M$ , and the density of the gas to be so small that the mutual action of the molecules could be neglected. The molecules were supposed to contain rapidly revolving charged particles.

If  $\theta$  is the angle which the magnetic axis of a molecule makes with

\* *Phil. Trans.*, 214 A, p. 109; 215 A, p. 79; 220 A, p. 247.

the direction of the external magnetic field  $H$ , then  $M\Sigma \cos\theta$  will be the resultant magnetic moment per unit volume or intensity of magnetization  $I$ , where the sign  $\Sigma$  indicates the summation of the values of  $\cos\theta$  for all the molecules in unit volume.

The problem is to find the value of  $\Sigma \cos\theta$  as a function of  $H$  and the absolute temperature  $T$ . In Chapter V on the quantum theory the equilibrium distribution of energy among a large number of similar systems, such as atoms, each of which is supposed to be only capable of having energies  $E_1, E_2, E_3 \dots$ , is considered. It is shown that the number having energies equal to  $E_n$  is proportional to  $e^{-E_n/kT}$ , where  $k$  is the gas constant for one molecule and  $T$  the absolute temperature. Hence if the energy is supposed to be capable of continuous variation the number having energies between  $E$  and  $E + dE$  must be proportional to  $e^{-E/kT} dE$ .

In a magnetic field  $H$  the potential energy of a molecule is  $MH(1 - \cos\theta)$ , where  $M$  is the magnetic moment and  $\theta$  the angle between  $H$  and the magnetic axis. Thus we may take for  $dN$ , the number of molecules in a unit volume for which  $\theta$  is between  $\theta$  and  $\theta + d\theta$ ,

$$dN = C e^{HM \cos\theta/kT} \sin\theta d\theta,$$

where  $C$  is a constant, since  $dN$  is proportional to  $e^{-E/kT} dE$ , and  $E = MH(1 - \cos\theta)$  so that  $dE = MH \sin\theta d\theta$ .

To determine  $C$  we have

$$N = C \int_0^\pi e^{HM \cos\theta/kT} \sin\theta d\theta.$$

Let  $\cos\theta = x$ , so that  $\sin\theta d\theta = -dx$ ,

$$\text{and} \quad N = C \int_{-1}^{+1} e^{HMx/kT} dx.$$

Hence

$$C = \frac{NHM}{kT(\epsilon^{HM/kT} - \epsilon^{-HM/kT})}.$$

The magnetic moment per unit volume or intensity of magnetization is therefore given by

$$\Sigma M \cos\theta = I = \int_0^\pi \frac{NHM^2 \epsilon^{HM \cos\theta/kT} \cos\theta \sin\theta d\theta}{kT(\epsilon^{HM/kT} - \epsilon^{-HM/kT})}$$

$$\text{or} \quad I = NM \left\{ \frac{\epsilon^{HM/kT} + \epsilon^{-HM/kT}}{\epsilon^{HM/kT} - \epsilon^{-HM/kT}} - \frac{kT}{HM} \right\}.$$

$$\text{Hence} \quad I = NM \left\{ \coth\left(\frac{HM}{kT}\right) - \frac{kT}{HM} \right\}.$$

According to this, when  $HM/kT$  is large, as at very low temperatures,

$$I = N \left\{ M - \frac{kT}{H} \right\};$$

and when  $HM/kT$  is small, as is usually the case,

$$I = \frac{NM^2H}{3kT},$$

so that the susceptibility is

$$\kappa = \frac{NM^2}{3kT}.$$

It was found by Curie that the susceptibility of most paramagnetic substances is approximately inversely as the absolute temperature, in agreement with Langevin's theory when  $HM/kT$  is small.

## 6. Extension of Langevin's Theory to Solids and Liquids.

Langevin's theory may be supposed to apply approximately to solids and liquids as well as to gases, provided they have very small susceptibilities, so that the field due to the atoms is small compared with the external field  $H$ .

The intensity of magnetization  $I$  is equal to the magnetic moment of the  $N$  molecules in unit volume so that, if  $\sigma$  denotes the magnetic moment of 1 gm.-molecule of any substance, then

$$\sigma = \frac{\mathcal{N}M^2H}{3kT},$$

where  $\mathcal{N}$  is the number of molecules in a gram-molecule of the substance. The saturation value of  $\sigma$  is  $\sigma_m = \mathcal{N}M$ , so that putting  $R = k\mathcal{N}$ , as usual, we get

$$\sigma = \frac{H\sigma_m^2}{3RT}.$$

This equation enables  $\sigma_m$  to be calculated when  $\sigma/H$  has been found at any temperature  $T$ .  $\sigma/H$  is called the molecular susceptibility and may be denoted by  $\chi$ , so that  $\chi = \sigma_m^2/3RT$ .

The product  $\chi T$  is nearly independent of  $T$  and  $H$  for paramagnetic substances at ordinary and higher temperatures. For example, for palladium Curie found it to be constant between 22° C. and 1370° C. However, it is found not to be constant at very low temperatures.

The following are the values of  $\chi$  found for anhydrous manganese sulphate at different temperatures.

Absolute Temperature.	$\chi \times 10^6$ .	$\chi T \times 10^6$ .
14.4	636	9,160
17.8	627	11,170
20.1	603	12,100
64.9	315	20,500
77.4	275	21,300
169.6	144	24,400
293.9	88	25,900

Thus the product  $\chi T$  becomes much smaller at very low temperatures.

The following are the values of  $\chi$  for ferrous sulphate,  $\text{FeSO}_4 + 7\text{H}_2\text{O}$ .

Absolute Temperature.	$\chi \times 10^6$ .	$\chi T \times 10^6$ .
14.7	756	11.100
20.3	571	11.600
64.6	191	12.300
77.3	160	12.400
292.3	42.4	12.400

In this case  $\chi T$  is much more nearly constant.

More elaborate theories of paramagnetism than Langevin's have been worked out by Gans and others. In these theories the mutual influence of the molecules is allowed for. The molecules are regarded as producing a field called the molecular field, which may be large compared with the external field  $H$ . The theory of this molecular field will be considered under ferromagnetism.

## 7. Modifications based on Quantum Theory.

Modifications of Langevin's theory based on the quantum theory have also been proposed by Keesom, Gans, Weyssenhoff, Reiche, and others. The general idea underlying these theories is similar to that on which the quantum theory of specific heats is based. The average energy of a gram-atom of a solid element on the classical theory is  $3RT$ , which makes the atomic heat equal to  $3R$  or about six calories. At low temperatures the atomic heats are less than six and become very small near the absolute zero of temperature, showing that the energy per gram-atom is less than  $3RT$ .

According to Debye's quantum theory of specific heats the energy per gram-atom of a solid element is given by

$$E = 9R \frac{T^4}{\psi^3} \int_0^{\psi/T} \frac{x^3 dx}{e^x - 1},$$

where  $\psi$  is a constant depending on the nature of the substance. When  $T$  is very large this formula gives  $E = 3RT$  in agreement with the classical theory.

Langevin's formula,

$$I = NM \left( \coth \frac{HM}{kT} - \frac{kT}{HM} \right),$$

with  $I_m = NM$ ,  $\sigma_m = \mathcal{N}M$ , and  $R = \mathcal{N}k$ , gives

$$\frac{I}{I_m} = \frac{\sigma}{\sigma_m} = \coth \left( \frac{H\sigma_m}{RT} \right) - \frac{RT}{H\sigma_m}.$$

We see that  $\sigma/\sigma_m$  is a function of  $H\sigma_m/RT$ , so that to obtain a quantum theory of paramagnetism we may replace  $RT$  in Langevin's formula

by  $E/3$  as given by Debye's formula. The constant  $\psi$  appropriate for calculating specific heats will be different from that to be used in the theory of paramagnetism, because paramagnetism depends only on the rotational oscillations of the molecules.

$$\text{This gives} \quad \frac{\sigma}{\sigma_m} = \coth\left(\frac{3H\sigma_m}{E}\right) - \frac{E}{3H\sigma_m},$$

$$\text{where} \quad E = 9R\frac{T^4}{\psi^3} \int_0^{\psi/T} \frac{x^3 dx}{e^x - 1}.$$

$$\text{At very low temperatures } E \text{ is small, so that } \coth\left(\frac{3H\sigma_m}{E}\right) = 1 \text{ and } E = \frac{3RT^4\pi^4}{5\psi^3}.$$

$$\text{Hence} \quad \sigma = \sigma_m - \frac{RT^4\pi^4}{5H\psi^3}.$$

If  $3H\sigma_m/E$  is small, we get

$$\frac{\sigma}{\sigma_m} = \frac{H\sigma_m}{E}, \text{ so that } \chi = \frac{\sigma_m^2}{E}.$$

By means of such quantum theories it is possible to explain the variation of  $\sigma$  with the temperature at low temperatures. The theoretical formulæ contain constants the values of which are selected so as to make the calculated results agree as well as possible with the experimental results. Under these circumstances, of course, the calculated results agree fairly well with those observed.

### 8. Ferromagnetism. Weiss's Theory.

In Langevin's theory of paramagnetism the magnetic molecules are supposed to oscillate in the external field  $H$  without any mutual action. Langevin's theory has been modified by Weiss, so as to be applicable to ferromagnetic bodies, by supposing that in such bodies there is a "molecular field" due to the mutual action of the magnetic molecules. This molecular field is assumed to be proportional to the intensity of magnetization  $I$  and directed parallel to it. Hence

$$H_m = \beta I,$$

where  $H_m$  denotes the molecular field strength and  $\beta$  is a constant depending on the nature of the substance. Weiss supposes that the magnetic molecules are arranged in sets or groups which may be small crystals of which magnetic metals like iron are composed, and that these groups are usually magnetized as strongly as possible even in an apparently unmagnetized piece of the metal.

In an unmagnetized bar the magnetic axes of the groups are sup-

posed directed at random, so that the total magnetic moment of any part of the bar containing a large number of groups is zero. An external field tends to make the magnetic axes of all the groups point along the direction of the external field, and when this has been accomplished the metal is magnetized to saturation.

Weiss supposes that Langevin's equation for a paramagnetic gas will apply to the groups of molecules in ferromagnetic bodies if  $H$  is replaced by the vector sum of  $H$  and  $\beta I$ , so that

$$I = NM \left\{ \coth \frac{M(H + \beta I)}{kT} - \frac{kT}{M(H + \beta I)} \right\}.$$

The intensity of magnetization  $I'$  of a group when  $H$  is small compared with  $\beta I$  is therefore given by

$$I' = NM \left\{ \coth \frac{\beta M I'}{kT} - \frac{kT}{\beta M I'} \right\}.$$

This equation gives  $I'$  as a function of the absolute temperature  $T$ . To solve it, let  $\frac{\beta M I'}{kT} = x$  and plot curves showing  $y = \coth x - \frac{1}{x}$

and  $y = \frac{I'}{NM} = \frac{kT}{\beta N M^2} x$  as functions of  $x$ . Let  $x'$  be the value of  $x$  at which these curves intersect (fig. 1), so that

$$I' = \frac{kT x'}{\beta M}$$

gives the intensity of magnetization of a group when the external field is negligible. If  $\sigma'$  denotes the magnetic moment per gram-molecule for this case, then  $\sigma' = I' \frac{\mathcal{N}}{N}$ ,

where  $\mathcal{N}$  is the number of molecules in a gram-molecule, so that since  $\mathcal{N}k = R$ ,

$$x' = \frac{\beta M N \sigma'}{RT}.$$

Now if  $\sigma_m$  denotes the saturation value of  $\sigma$ , i.e. the value of the magnetic moment per gram-molecule when  $H$  is very great, then

$$\sigma_m = MN \frac{m}{D},$$

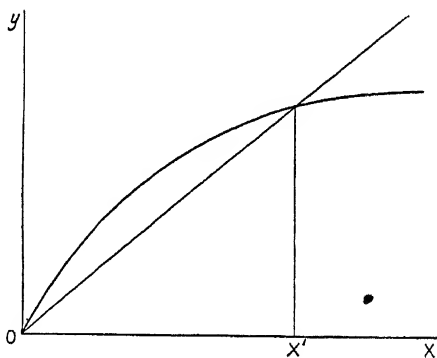


Fig. 1

where  $m$  is the molecular weight of the magnetic molecules and  $D$  the density of the substance, so that

$$x' = \frac{\beta \sigma' \sigma_m D}{RTm}.$$

Hence

$$\sigma' = \frac{mRTx'}{\beta D \sigma_m}.$$

As the temperature is raised the slope of the line  $y = \frac{kT}{\beta NM^2} x$  increases, so that the value of  $x'$  diminishes until a temperature is reached at which  $x'$  and therefore  $\sigma'$  become zero. This is the critical temperature at which the ferromagnetic properties disappear. When  $x' = 0$ , the curves  $y = \coth x - \frac{1}{x}$  and  $y = \frac{kT}{\beta NM^2} x$  touch each other at the origin, so that  $dy/dx$  has the same value for both curves.

When  $x$  is very small the equation

$$y = \coth x - \frac{1}{x}$$

reduces to  $y = \frac{x}{3}$ , so that  $\frac{dy}{dx} = \frac{1}{3}$ , and so at the critical temperature  $T_c$  we have

$$\frac{kT_c}{\beta NM^2} = \frac{1}{3},$$

or

$$T_c = \frac{1}{3} \frac{\beta NM^2}{k} = \frac{\beta D \sigma_m^2}{3Rm}.$$

With

$$\sigma' = \frac{mRTx'}{\beta D \sigma_m}$$

this gives

$$\frac{T}{T_c} = \frac{3\sigma'}{x' \sigma_m}.$$

Now let

$$\frac{T}{T_c} = \theta, \quad \text{and} \quad \frac{\sigma'}{\sigma_m} = \phi,$$

so that

$$\theta = \frac{3\phi}{x'}.$$

We have also

$$\phi = \frac{\sigma'}{\sigma_m} = \coth x' - \frac{1}{x'}.$$

The equations

$$\theta = \frac{3\phi}{x'} \quad \text{and} \quad \phi = \coth x' - \frac{1}{x'}$$

give

$$\phi = \coth \frac{3\phi}{\theta} - \frac{\theta}{3\phi}.$$

This equation contains nothing depending on the nature of the particular substance considered, so that if the temperature is expressed as a fraction of the critical temperature, and the intensity of mag-

netization of the molecular groups, in zero field, as a fraction of its saturation value, then the relation between these quantities so expressed should be the same for all substances.

Now it is found that the constant  $\beta$  is very large, so that the molecular field is usually very large compared with the external field  $H$ . The intensity of magnetization of the molecular groups is therefore practically uninfluenced by the external field. It is therefore equal to  $\sigma'$  whatever may be the value of the external field, except near the critical temperature, when  $\sigma'$  becomes very small. The external field therefore merely acts by lining up the molecules in the groups so that they all point along the direction of the external field. According to this the saturation intensity of magnetization observed in strong external fields is not  $\sigma_m$  but  $\sigma'$ . Thus measurements of  $\sigma$  in strong fields give  $\sigma'$ , so that it is possible to test the equation  $\phi = \coth \frac{3\phi}{\theta} - \frac{\theta}{3\phi}$  experimentally.

The curve in fig. 2 shows the relation between  $\theta$  and  $\phi$  given by this equation, and the points marked represent the experimental results found by Weiss for magnetite in a field of 8300 gauss. The agreement is good, but in the cases of iron, nickel, and cobalt the differences between the theory and the experimental results are greater than with magnetite. Thus the theory gives a fairly satisfactory explanation of the variation of the saturation intensity of magnetization of ferromagnetic substances with the

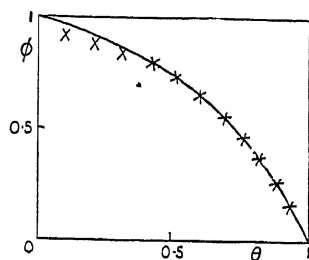


Fig. 2

temperature.

The value of the constant  $\beta$  has been found by making measurements of the intensity of magnetization near the critical temperature, where  $\beta I$  is not too large compared with  $H$  for the effects due to  $H$  to be appreciable. Near the critical temperature  $\frac{M(H + \beta I)}{kT}$  is small, so

that we have

$$I = \frac{NM^2(H + \beta I)}{3kT}.$$

With  $T_c = \frac{1}{3} \frac{\beta NM^2}{k}$ , this gives  $\frac{I\beta}{H} = \frac{T_c}{T - T_c}$ .

It was found by Curie that near the critical temperature the susceptibility  $\kappa = I/H$  for ferromagnetic substances is inversely proportional to  $T - T_c$  in agreement with this equation, so that  $\beta$  can be calculated from such measurements of  $\kappa$ . In this way it is found that  $\beta$  has the following values:

	$\beta$ .	$\beta I'$ .
Iron .. ..	3,850	6,560,000
Nickel .. ..	12,700	6,350,000
Magnetite ..	33,200	14,300,000
Cobalt .. ..	6,180	8,870,000

The values given under  $\beta I'$  are the values of the molecular field in the groups of molecules at the ordinary temperature. These values are much larger than the external fields which have been used, which are seldom greater than 30,000 gauss, so that it is clear that the external fields can make no appreciable difference to  $I'$  at ordinary temperatures.

It is difficult to see how such enormous fields can be produced by the molecular magnets, and if such fields exist in the molecular groups it is difficult to see how the external field can line them up.

The hypothesis of the molecular field has, however, been of great value, and it certainly enables many important magnetic phenomena to be explained.

## 9. Weiss's Explanation of Hysteresis.

To explain the observed relations between the intensity of magnetization and the external field, and in particular the phenomena of hysteresis, Weiss suggested that the molecular groups which are supposed to be magnetized in random directions with intensity  $I'$  have their directions of magnetization reversed by a comparatively weak field in a direction opposite to the direction of magnetization. Let  $H_c$  be the field strength required to reverse  $I'$  in a group, and let  $\theta$  be the angle between the magnetic axis of a group and the external field  $H$ . Then we suppose all directions of the magnetic axes of the groups equally probable, so that half the groups will not be affected by the external field however strong it becomes. The other half will not be affected until  $H \cos \theta$  is greater than  $H_c$ , and then the directions of their axes will be reversed.

All groups for which  $\theta$  is between  $\pi - \bar{\theta}$  and  $\pi$  will be reversed, where  $H \cos \bar{\theta} = H_c$ , and the resultant intensity of magnetization will be twice that due to these groups. Hence

$$I = -I' \int_{\pi-\bar{\theta}}^{\pi} \sin \theta \cos \theta d\theta = \frac{I'}{2} \sin^2 \bar{\theta},$$

$$\text{so that } I = \frac{I'}{2} \left\{ 1 - \left( \frac{H_c}{H} \right)^2 \right\}.$$

If  $H$  is increased from 0 up to a certain value and then diminished,  $I$  will remain constant at the value given by this equation as  $H$  is diminished, until  $H = -H_c$ , when it will begin to diminish with  $H$ . If the maximum value of  $H$  is much greater than  $H_c$  the maximum value of  $I$  will be  $I'/2$ , and when  $H$  is diminished and reversed then  $I$ , when  $-H$  is greater than  $H_c$ , will be given by

$$I = -\frac{I'}{2} \left\{ 1 - 2 \left( \frac{H_c}{H} \right)^2 \right\}.$$

In this case  $I$  will be zero when  $H = -\sqrt{2}H_c$ . In this way an hysteresis

curve is obtained rather like the curves actually obtained with iron. Weiss also supposed that the molecular groups were magnetized slightly by the field  $H$  in directions perpendicular to their axes, and so obtained theoretical curves rather more like those found experimentally. However, it is clear that this theory of hysteresis is inadequate.

## 10. Other Theories of Ferromagnetism.

A theory of ferromagnetism similar to that of Weiss, but more elaborate, has been worked out by Gans. He assumes that in addition to the external field  $H$  and the molecular field  $\beta I$  there is a third field  $\frac{1}{3}\pi I$ . Gans supposes that the molecular field assumes different directions and that all directions are equally probable. This means that the molecular groups are continually changing the directions of their axes. The results of Gans's theory are very similar to those of the theory of Weiss.

A theory of the relation between  $H$  and  $I$  has been developed by Honda and Okubo. This theory is an elaboration of the classical theory of Ewing. They consider first a Ewing model consisting of nine magnets. These magnets are supposed placed on a square, one at each corner, one in the middle of each side, and one in the middle of the square. The magnets are supposed free to turn in any direction except in so far as they are influenced by the external field and their own fields.

Honda and Okubo worked out the relation between the moment of such a group and the external field, and they then went on to consider the case of a large number of such groups orientated at random, which may be supposed to represent a ferromagnetic substance. They show that the relation between  $I$  and  $H$  for such a set of groups is very similar to that observed in soft iron. This agrees with the experimental results obtained long ago by Ewing, who showed that the magnetic properties of iron could be imitated by means of a large number of freely suspended small magnets equally spaced over a plane area.

Ewing has recently developed a modification of his original theory. A group of small magnets equally spaced on a lattice has too much stability to represent the behaviour of soft iron in weak fields. When the magnets are lined up so that each north pole is opposite the south pole of the next magnet it requires a very strong field to reverse the direction of the group. Ewing therefore suggests a model of a ferromagnetic atom, which consists of a small magnet free to rotate in any direction surrounded by eight fixed magnets with their centres on the corners of a cube and their axes all directed towards the small magnet at the centre of the cube. He shows that this arrangement gives the movable magnet the desired small stability, since if its south pole is near one of the fixed north poles then its north pole must be near another fixed north pole. It is then very easily deflected from one such position to another. In a group of such atoms the mutual action of the movable magnets will be added to the action of the fixed magnets on the movable ones. Ewing considers that such a model may resemble an iron atom, and that it gives a better account of the relations between field and magnetization than his original model of equally spaced freely rotatable magnets.

## 11. Magnetic Properties of Crystals.

The magnetic properties of several ferromagnetic crystals have been carefully investigated by Weiss and others. The case of "normal" pyrrhotite,  $\text{FeS}$ , from Brazil, will be considered here. This crystal is much more easily magnetized in one direction than in any other.

Taking this direction as the  $x$  axis, then in the  $yz$  plane there is a

direction, which will be taken as the  $y$  axis, in which the crystal is much more easily magnetized than in the perpendicular direction. A field of about 15 gauss along the  $x$  axis is sufficient to reverse the direction of magnetization, and a field about double this produces saturation. The hysteresis loop for fields along the  $x$  axis resembles that obtained with very pure iron. Along the  $y$  axis a field of about 730 gauss is required to reverse the magnetization and about 12,000 gauss to produce approximate saturation. The crystal cannot be magnetized to any extent along the  $z$  axis.

The intensity of magnetization due to a field of 12,000 gauss or more in the  $xy$  plane, or the magnetic plane as it is called, is the same in all directions and is in the direction of the field. It is equal to 47. With weaker fields the intensity is 47 when the field is along the  $x$  axis but is less along  $y$ , and for intermediate directions has intermediate values with the direction of the magnetization nearer to  $Ox$  than the direction of the field.

If  $H_x, H_y, H_z$  are the components of the external field and  $I_x, I_y, I_z$  those of the intensity of magnetization, then Weiss supposes that the total field, or resultant of the external field and the molecular field, has components  $H_x + \beta_x I_x, H_y + \beta_y I_y, H_z + \beta_z I_z$ , where  $\beta_x, \beta_y, \beta_z$  are constants which are unequal. Weiss assumes that the resultant intensity of magnetization is in the same direction as the resultant field, so that

$$\frac{H_x + \beta_x I_x}{I_x} = \frac{H_y + \beta_y I_y}{I_y} = \frac{H_z + \beta_z I_z}{I_z} = \alpha.$$

The quantity  $\alpha$  is nearly constant for weak fields. We have therefore

$$I_x = \frac{H_x}{\alpha - \beta_x}, \quad I_y = \frac{H_y}{\alpha - \beta_y}, \quad I_z = \frac{H_z}{\alpha - \beta_z}.$$

If  $H$  lies in the  $xy$  plane and makes an angle  $\theta$  with the  $x$  axis, then

$$\frac{H \cos \theta + \beta_x I \cos \phi}{I \cos \phi} = \frac{H \sin \theta + \beta_y I \sin \phi}{I \sin \phi},$$

where  $\phi$  is the angle between  $I$  and the  $x$  axis. Hence

$$I = \frac{H \sin(\theta - \phi)}{(\beta_x - \beta_y) \sin \phi \cos \phi}.$$

The assumption that the intensity  $I$  is along the direction of the resultant field is equivalent to supposing that the axes of the molecular magnets are symmetrically arranged about this direction.

A similar expression to the above may easily be obtained for fields in the  $xz$  plane. It is found that the experimental results agree approximately with these formulæ. The value of  $\beta_x - \beta_y$  is found to be 153

and that of  $\beta_x - \beta_z$  is 3200. The saturation value of  $I$  is 47. The assumption of the existence of a molecular field enormously greater than the intensity of magnetization is difficult to justify, but it appears to give results in agreement with the facts. This field of course may represent forces on the molecules of other than magnetic origin.

The magnetic properties of magnetite,  $\text{Fe}_3\text{O}_4$ , hematite,  $\text{Fe}_2\text{O}_3$ , and of iron crystals have also been studied. They are somewhat more complicated than those of pyrrhotite.

## 12. Magnetization and Rotation.

Since the magnetic moment of magnetic atoms or molecules is supposed to be due to electrons describing orbits in them, or possibly to spinning electrons, it is clear that a magnetic molecule must have a moment of momentum about its magnetic axis. It was shown in section 3 that the angular momentum due to free electrons is equal per unit volume to  $2I \frac{m}{e}$ , where  $I$  is the intensity of magnetization,  $m$  the mass, and  $e$  the charge of an electron. It can easily be shown that this result is true for magnetization due to electrons describing orbits of any kind. According to this, when a bar is magnetized it acquires internal angular momentum  $2I \frac{m}{e}$  per unit volume. Since the total angular momentum of the bar must remain constant provided there is no couple acting on it, we should expect the bar to be set rotating about its magnetic axis with angular momentum equal and opposite to the internal angular momentum. This was pointed out by O. W. Richardson in 1914. The effect has since been detected experimentally by Einstein and Haas, and has been carefully measured by several observers. It is found that the observed effect is equal to  $I \frac{m}{e}$  instead of  $2I \frac{m}{e}$  as predicted by Richardson. The converse of Richardson's effect was discovered by Barnett in 1915. He showed that when a bar of iron is set rotating about its axis it becomes slightly magnetized.

## 13. Measurement of Magnetic Moment of Atoms.

Stern and Gerlach have introduced a new method of measuring the magnetic properties of atoms, and this method has already given very interesting results. Vapour of the substance to be investigated is allowed to pass through a narrow slit into a vacuum. The molecules move along straight lines in the vacuum, so forming a diverging beam. By means of a second slit parallel to the first a narrow nearly parallel beam is obtained. This beam is passed between the poles of a magnet which gives a strong field perpendicular to the plane containing the slits. The field strength is made to vary as rapidly as possible along

the direction of the field. After passing through this field the molecules fall on a screen on which they are condensed and make a visible mark. In this way the deflection of the molecules by the magnetic field can be measured.

Take the  $x$  axis along the original direction of the stream of molecules from the second slit and the  $y$  axis along the direction of the magnetic field. If  $M$  denotes the magnetic moment of a molecule then the force on it due to the field is  $M_y \frac{\partial H}{\partial y}$ , where  $M_y$  denotes the  $y$  component of  $M$ . The deflection due to the field is therefore given by

$$y = \frac{1}{2} \frac{M_y}{A} \frac{\partial H}{\partial y} \frac{l^2}{v^2},$$

where  $A$  is the mass of the molecule,  $v$  its velocity, and  $l$  the distance from the second slit to the screen.

The average value of  $v^2$  can be calculated from the temperature of the vapour, so that  $M_y$  can be deduced from the deflection.

In this way it was found that the atoms of copper, silver, and gold have magnetic moments of about  $9 \times 10^{-21}$ . Some of the atoms were deflected in the direction of the magnetic field as though they had moments  $+9 \times 10^{-21}$ , and others were deflected equally in the opposite direction as though they had moments  $-9 \times 10^{-21}$ .

According to Bohr's quantum theory, the plane of the orbit of the outer electron must be perpendicular to the magnetic field and the electron may revolve round the orbit in either direction. The angular momentum of this electron according to the quantum theory is  $\hbar/2\pi$ , where  $\hbar$  is Planck's constant. Hence for a circular orbit of radius  $a$  we have  $mva = \pm \hbar/2\pi$ , where  $v$  is the orbital velocity. The magnetic moment is the frequency  $v/2\pi a$  multiplied by the electronic charge  $e$  and the area of the orbit, so that  $M = \frac{v}{2\pi a} e\pi a^2 = \frac{vea}{2}$ . But  $va = \pm \hbar/2\pi m$ , so that  $M = \pm \frac{\hbar e}{4\pi m}$ , according to the quantum theory. We have  $\hbar = 6.55 \times 10^{-27}$  and  $e/m = 1.77 \times 10^9$ , so that  $M = 9.21 \times 10^{-21}$ , which agrees well with the values found experimentally. The magnetic moment  $\hbar e/4\pi m$  is sometimes called the moment of a Bohr magneton.

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## CHAPTER III

# Thermionics

### 1. Experimental Phenomena.

When a body charged with electricity is heated to a sufficiently high temperature the electricity leaks away and the body becomes discharged. The branch of physics dealing with such phenomena was called *thermionics* by O. W. Richardson, to whom our knowledge of it is largely due.

A simple form of apparatus for studying thermionics is shown in fig. 1. A loop of wire AB of tungsten, platinum, or any other metal having a high melting-point is supported by two platinum wires EF sealed into a glass bulb as shown. The loop is surrounded by a metal cylinder CD supported by wires sealed through the glass at the lower end of the bulb. A tube T connects the bulb to apparatus for producing a good vacuum in it.

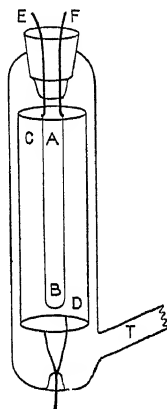


Fig. 1

The loop can be heated by passing a current through it, and its temperature can be found from its resistance or by means of an optical pyrometer.

The cylinder is connected through a galvanometer to one terminal of a battery by means of which any desired potential difference can be maintained between the loop and the cylinder. The other terminal of the battery is connected to one of the wires E and F. If a good vacuum is maintained in the bulb, that is if the gas pressure in it is kept below say  $10^{-4}$  of a millimetre of mercury, then on raising the temperature of the loop it is found that a current is indicated by the galvanometer when the loop is negatively charged, but that there is no current when it is positively charged.

Negative electricity escapes from the wire to the cylinder when the wire is hot enough, but not positive electricity. The galvanometer indicates a negative current flowing into the cylinder when the cylinder is at a higher potential than the wire.

The current obtained depends on the temperature and on the

potential difference. If the temperature is kept constant, and the potential difference gradually increased from zero, then the current at first increases rapidly with the potential difference but soon attains a constant maximum value. This maximum current is called the *saturation current*. The potential required to produce the saturation current is greater when the saturation current is large than when it is small. When the temperature is raised the saturation current increases rapidly with it.

## 2. O. W. Richardson's Theory of the Thermionic Current.

The negative electricity which escapes from hot metals in a good vacuum consists of negative electrons. That this is the case was first shown by J. J. Thomson, who measured the ratio of the charge  $e$  to the mass  $m$  of the escaping electricity, and found it equal to about  $10^7$  electromagnetic units per gram, which is about the value of  $e/m$  for negative electrons.

The electrical conductivity of metals is supposed to be due to the presence in them of negative electrons which are free to move about inside the metal. It is natural to suppose that the electrons which escape from the metal at high temperatures are some of these free electrons.

The free electrons were at one time supposed to have the same average kinetic energy as the molecules of a gas at the same temperature, and a theory of the negative thermionic current based on this supposition was worked out by O. W. Richardson.

This classical theory is not now regarded as correct, but a brief account will be given of it here because of its historical interest. The more recent theories based on thermodynamics and the quantum theory will then be discussed.

We suppose that a metal contains  $n$  free electrons per unit volume and that these electrons move about inside the metal like the molecules of a gas, colliding with each other and with the metallic atoms. The average kinetic energy of the free electrons we suppose is equal to  $\alpha T$ , where  $T$  is the absolute temperature of the metal and  $\alpha$  is a constant.

Take axes  $x, y, z$  in the metal and let  $u, v, w$  be the components of the velocity  $V$  of an electron parallel to these axes. Then we suppose that the distribution of the velocity components among the electrons is given by Maxwell's law for gas molecules, that is, if  $dn$  denotes the number in unit volume having velocity components  $u$  between  $u$  and  $u + du$ , then

$$dn = nA\varepsilon^{-qu^2} du,$$

where  $A$  and  $q$  are constants at any given temperature.

We have 
$$\int_{-\infty}^{+\infty} dn = \int_{-\infty}^{+\infty} nA\varepsilon^{-qu^2} du = n,$$

and 
$$3 \int_{-\infty}^{+\infty} \frac{1}{2} mu^2 dn = \int_{-\infty}^{+\infty} nA \frac{3}{2} mu^2 \varepsilon^{-qu^2} du = n\alpha T.$$

But 
$$\int_{-\infty}^{+\infty} \epsilon^{-qu^2} du = \sqrt{\frac{\pi}{q}}, \text{ and } \int_{-\infty}^{+\infty} u^2 \epsilon^{-qu^2} du = \frac{\sqrt{\pi}}{2q^{3/2}},$$

so that 
$$A \sqrt{\frac{\pi}{q}} = 1, \text{ and } \frac{3mA \sqrt{\pi}}{4q^{3/2}} = \alpha T,$$

which give 
$$A = \sqrt{\frac{3m}{4\pi\alpha T}}, \text{ and } q = \frac{3m}{4\alpha T}.$$

For gas at pressure  $p$  containing  $n$  molecules per unit volume we have  $p = \frac{1}{3}nm\bar{V}^2$ , where  $m$  is the mass of one molecule and  $\bar{V}^2$  the average value of the square of the velocities of the molecules.

### 3. The Thermionic Work Function.

Richardson assumed that before an electron can escape from the metal it has to do a definite amount of work at the surface layer. Let us denote this work by  $\phi e$ , where  $e$  is the charge on one electron, so that  $\phi$  is the potential difference corresponding to the work in question. According to this, if  $\frac{1}{2}mV_1^2$  is the kinetic energy of an electron in the metal before escaping, and  $\frac{1}{2}mV_2^2$  that after escaping, then

$$\frac{1}{2}m(V_1^2 - V_2^2) = \phi e.$$

The number of electrons which enter the surface layer per unit area in unit time is

$$\int u dn = \int_0^\infty nA \epsilon^{-qu^2} u du,$$

where  $u$  is the velocity component towards the surface, which we take to be perpendicular to the  $x$  axis. Of these only those for which  $\frac{1}{2}mu^2$  is greater than  $\phi e$  can escape. The number escaping per unit area in unit time is therefore

$$\int_{\sqrt{\frac{2\phi e}{m}}}^\infty nA \epsilon^{-qu^2} u du = \frac{n \epsilon^{-2\phi eq/m}}{2\sqrt{\pi q}},$$

since  $A = \sqrt{\frac{q}{\pi}}$ .

The thermionic current density  $i$  is therefore given by

$$i = \frac{ne \epsilon^{-2\phi eq/m}}{2\sqrt{\pi q}}.$$

Substituting  $3m/4\alpha T$  for  $q$ , this becomes

$$i = ne \sqrt{\frac{\alpha T}{3\pi m}} \epsilon^{-\frac{\phi e}{\alpha T}}.$$

If we put

$$a = ne \sqrt{\frac{\alpha}{3\pi m}} \text{ and } b = \frac{\phi e}{\alpha},$$

we get

$$i = a T^{1/2} \epsilon^{-b/T}.$$

It is found that the saturation current density varies with the temperature approximately in accordance with this equation. The factor  $\epsilon^{-b/T}$  increases very rapidly with  $T$ , so that the effect of the factor  $T^{1/2}$  is scarcely appreciable. However, recent very exact measurements of  $i$  seem to show that the power of  $T$  should be about 2 instead of  $\frac{1}{2}$ .

The following table gives some values of the thermionic work function  $\phi$  expressed in volts:

Tungsten	..	..	4.52
Platinum	..	..	4.4
Iron	..	..	3.7
Aluminium	..	..	3.0
Sodium	..	..	1.8

The work function  $\phi$  is closely related to contact potential difference. Consider a condenser consisting of two parallel plates made of different metals, and let the two plates be connected together by a wire of the same metal as one of the plates is made of.

Suppose an electron is taken from a point inside one of the plates through the wire into the other plate and then across the space between the plates back to its original position. The total work required to take the electron around this path with negligible velocity must be zero. If we neglect the small potential difference at the junction of the two metals, the work required is

$$e(\phi_1 - \phi_2) + e(V_2 - V_1) = 0,$$

where  $V_2$  is the potential just outside one metal and  $V_1$  that just outside the other. But  $V_2 - V_1$  is the contact potential difference, so that the difference between the thermionic work functions for two metals is nearly equal to the contact potential difference between them. Cf. Chap. IV, section 2.

#### 4. Objections to Classical Theory.

When light falls on the surface of a metal it may cause electrons to escape. This is the photoelectric effect (see Chap. IV). The maximum kinetic energy of the electrons which escape is given by Einstein's equation,

$$\frac{1}{2}mv^2 = h\nu - w,$$

where  $\nu$  is the frequency of the light,  $h$  is Planck's constant, and  $w$  a constant depending on the nature of the metal.  $h\nu$  is supposed to be the energy of one quantum of the light of frequency  $\nu$ , so that if an electron in the metal absorbs a quantum then it gets energy  $h\nu$ .

It is found that  $w$  is equal to  $\phi e$ , the work which an electron must do in escaping through the surface layer. This seems to require that the electrons set free by the light should have no appreciable kinetic energy in the metal when they absorb the quantum of energy  $h\nu$ . This does not agree with Richardson's assumption that the electrons have the same average kinetic energy as gas molecules.

The specific heat of electricity in metals is also much smaller than it should be if the electrons carrying the current have the same energy as gas molecules. Also it is found that the specific heats of metals can be explained without attributing any thermal kinetic energy to the

electrons in them. For these reasons the idea that the average kinetic energy of the free electrons in metals is equal to that of gas molecules has been abandoned. It seems probable that the electrons in metals are in stationary states of the quantum theory, so that so long as they do not change from one such state to another their energy is independent of the temperature of the metal.

### 5. Thermodynamical Theory of Thermionic Emission.

If we assume that the escaping electrons form a monatomic gas or vapour which can be in equilibrium with the metal when at a definite pressure, then we can apply the thermodynamical theory of evaporation to the problem and so obtain a theory of thermionic emission without making assumptions as to the state of the electrons inside the metal. A theory of this kind was first proposed by the writer and has since been developed by O. W. Richardson and others.

Let us first consider the latent heat of evaporation of one mol of electrons. Let the potential difference between a point just outside the metal and a point inside it be  $\phi$ . Then if  $\mathcal{N}$  is the number of molecules in one mol of any gas and  $e$  the charge on one electron, the work required when one mol of electrons escapes from the metal is  $\mathcal{N}e\phi$ .

As we have seen, the electrons in the metal have little or no kinetic energy, whereas in the gaseous state outside they have the same average kinetic energy as the molecules of a gas. For any gas (p. 49) we have  $p = \frac{1}{3}mn\bar{V}^2$  or  $p = \frac{2}{3}naT$ , since we have taken (p. 48)  $aT$  to be the average kinetic energy of the free electrons. Thus if we write  $k = \frac{2}{3}a$  we have  $p = nkT$ , where  $k$  is the gas constant for one molecule. The kinetic energy of the molecules in one mol of a monatomic gas is therefore equal to  $\frac{3}{2}\mathcal{N}kT = \frac{3}{2}RT$ , where  $R = \mathcal{N}k$  is the gas constant for one mol of any gas.

Also when the electrons escape as a gas at a pressure  $p$ , work  $pV$  has to be done against the external pressure  $P$ . But  $pV = RT$ , so that the total heat energy required to keep the temperature constant when one mol of electrons escape is

$$\mathcal{N}e\phi + \frac{5}{2}RT.$$

If then  $L$  denotes the heat of evaporation at the temperature  $T$ , and  $L_0 = \mathcal{N}e\phi_0$  that at the absolute zero of temperature,

$$L = L_0 + \mathcal{N}e(\phi - \phi_0) + \frac{5}{2}RT,$$

where  $\phi_0$  is the value of  $\phi$  at  $T = 0$ .

Contact difference of potential is found to be practically independent of the temperature, so that it is probable that  $\phi - \phi_0$  is very small, and we have approximately

$$L = L_0 + \frac{5}{2}RT.$$

It is only allowable to regard the electron gas as a perfect gas when its pressure and its volume are very small so that the electrical forces due to the charges on the electrons can be neglected.

This introduces no difficulty, because we need not suppose that more than an infinitesimal number of electrons are outside the metal at any time. In the chapter on the quantum theory (Chap. V, section 8) it is shown that the vapour pressure of a liquid or solid which gives off a monatomic vapour is given approximately by the equation:

$$\log p = -\frac{L}{RT} + \frac{5}{2} \log T + \frac{5}{2} + \log \left\{ \left( \frac{2\pi}{h^2} \right)^{3/2} k^{5/2} \right\} + \frac{3}{2} \log m,$$

where  $h$  is Planck's constant,  $L$  the heat of evaporation per mol at the temperature  $T$ , and  $m$  the mass of one molecule of the vapour. Since the electrons form a monatomic gas outside the metal we may take the pressure  $p$  at which they are in equilibrium with the metal to be given by this equation. Hence, putting  $L = L_0 + \frac{5}{2}RT$ , we get

$$\log p = -\frac{L_0}{RT} + \frac{5}{2} \log T + \log \left\{ \left( \frac{2\pi}{h^2} \right)^{3/2} k^{5/2} \right\} + \frac{3}{2} \log m.$$

The pressure  $p$  is the very small pressure of the electron gas which is in equilibrium with the metal. The relation between  $p$  and the saturation current density  $i$  may be determined if we assume that the electrons in the gas which collide with the metal surface are all absorbed and do not rebound. This is believed to be approximately true. Assuming this, we have the result that when the gas is in equilibrium with the metal the number of electrons which escape from the metal is equal to the number which collide with it.

We suppose that the electron gas can be regarded as a perfect gas, and that the average kinetic energy of the electrons in it is the same as for any other gas at the same temperature.

The number of electrons colliding with unit area of the metal surface in unit time is therefore given by the same expression as the number entering the surface layer from the metal on Richardson's classical theory, provided that  $n$  is now taken to be the number of electrons per unit volume in the electron gas instead of inside the metal. The number striking unit area in unit time is therefore (section 2)

$$\int_0^\infty nA e^{-u^2} u du = \frac{n}{2\sqrt{\pi q}}.$$

But  $q = \frac{3m}{4\alpha T}$  (p. 49),  $p = nkT$ , and  $\alpha = \frac{3}{2}k$  (p. 51), so that

$$\frac{n}{2\sqrt{\pi q}} = \frac{p\sqrt{2kT}}{2kT\sqrt{\pi m}} = \frac{p}{\sqrt{2\pi mkT}}.$$

The thermionic saturation current density  $i$  is therefore given by

$$i = \frac{pe}{\sqrt{2\pi mkT}},$$

where  $e$  is the charge on one electron.

Eliminating  $p$  from the equation for  $\log p$  and this expression for  $i$  we get

$$i = \frac{2\pi emk^2}{h^3} T^2 e^{-L_0/RT}.$$

This theoretical expression for  $i$  is based on the thermodynamical theory of evaporation and on the quantum theory. It was first obtained by Dushman.

The values of  $m, k, h$ , and  $R$  are all known with considerable accuracy, and the only quantity in the expression for  $i$  which depends on the properties of the particular metal emitting the electrons is  $L_0 = \mathcal{N}e\phi_0$ .

If we put  $\frac{2\pi emk^2}{h^3} = A$  and  $L_0/R = b$ , the equation for  $i$  becomes  $i = AT^2 e^{-b/T}$ , which gives  $b = T \log \left( \frac{AT^2}{i} \right)$ . The following tables give the values of  $i$  for tungsten and platinum at different temperatures measured by Davisson, Germer, and Schlichter, and the values of  $b$  calculated from them by means of this equation.

Temperature.	Current.	$b$
TUNGSTEN		
1 = 10 <sup>-2</sup> amp. per sq. cm.		
1935.5	0.0934	51,890
1986.5	0.1973	51,880
2036.0	0.3967	51,860
2077.5	0.6784	51,880
2086.5	0.7656	51,900
2102.0	0.9363	51,840
2131.5	1.362	51,840
2134.5	1.419	51,870
2158.0	1.902	51,820
2182.0	2.538	51,810
2204.0	3.269	51,820
2231.0	4.405	51,820
2235.0	4.606	51,870
2271.5	6.875	51,880
2280.0	7.394	51,920
2306.0	9.792	51,900

Temperature.	Current.	$b$
PLATINUM		
$1 = 10^{-8}$ amp. per sq. cm.		
1211.0	0.42	49,300
1243.0	1.35	49,200
1275.0	4.17	49,100
1307.0	9.9	49,300
1339.0	27.5	49,200
1371.0	76.0	49,000
1403.0	168.0	49,100
1435.0	400.0	49,100
1467.0	785.0	49,300
1499.0	1430.0	49,500

It will be seen that the values of  $b$  are all very nearly equal, so that the equation  $i = AT^2 e^{-b/T}$  represents the experimental results very well indeed.

## 6. Space Charge Effect.

When the potential difference used is not sufficient to produce the saturation value of the current, the current varies with the potential difference. If, for example, the current obtained is only one-half of the saturation current, then half the electrons emitted by the wire get across to the surrounding electrode and half return to the wire. The negative charge on the electrons in the space around the hot wire modifies the electric field and actually reverses it near the wire, so that the electrons coming out of the wire have to move a certain distance in a field which tends to stop them. Only those which have sufficient kinetic energy are able to get through this region in which the field is reversed, and the rest are driven back into the wire. This is called the space charge effect. To simplify the theory we will consider the case of a large plane area emitting electrons to a large plane electrode at a distance  $d$  from the emitting plane. Let  $x$  denote the distance of a point from the emitting plane, and  $V$  the potential difference between the point and the emitting plane. Then we have

$$\frac{\partial^2 V}{\partial x^2} = -4\pi ne,$$

where  $ne$  is the density of the space charge,  $n$  being the number of electrons per unit volume and  $e$  the charge on one electron. Let the region in which the field is reversed be of thickness  $t$ . When  $x$  is greater than  $t$  all the electrons present are moving away from the emitting plane, so that if their average velocity is  $v$  we have  $i = nev$ .

The velocity of an electron will be given by the equation

$Ve = \frac{1}{2}m(v_0^2 - v^2)$ , where  $v_0$  is the initial velocity of emission from the hot plane. The velocity components parallel to the emitting plane make no difference, so we disregard them and take  $v$  and  $v_0$  to be perpendicular to the electrodes. In the region of reversed field  $V$  is negative so that  $v$  is less than  $v_0$ . Let the minimum potential at  $x = t$  be  $V_1$ . Then if  $v_0$  is not greater than  $\sqrt{\frac{2V_1e}{m}}$  the electron will not get across and will return to the emitting plane.

The average velocity of the electrons at  $x = t$  will be the same as the average velocity of emission  $v_0$  at  $x = 0$ , and we shall suppose that this is small compared with  $v$ . The average velocity will then be given approximately by  $-Ve = \frac{1}{2}mv^2$ . Eliminating  $n$  and  $v$  from the three equations

$$\frac{\partial^2 V}{\partial x^2} = -4\pi ne,$$

$$i = nev,$$

$$Ve = -\frac{1}{2}mv^2,$$

we get

$$\frac{\partial^2 V}{\partial x^2} = -2\pi i \sqrt{\frac{-2m}{Ve}}.$$

Integrating this equation we get

$$\left(\frac{\partial V}{\partial x}\right)^2 = -8\pi i \sqrt{\frac{-2mV}{e}} + \text{constant}.$$

To determine the constant, we have  $\frac{\partial V}{\partial x} = 0$  at  $x = t$ , since  $V$  is a minimum at this point. We shall suppose that  $t$  is small, so that approximately  $\frac{\partial V}{\partial x} = 0$  when  $x = 0$  and  $V = 0$ , and the constant is zero.

Hence

$$\left(\frac{\partial V}{\partial x}\right)^2 = -8\pi i \sqrt{\frac{-2mV}{e}}.$$

Integrating this and putting  $V = 0$  at  $x = 0$ ,

we get

$$i = -\frac{1}{9\pi} \sqrt{\frac{-2e}{m}} \frac{V^{3/2}}{x^2}.$$

This gives

$$i = -2.33 \times 10^{-6} \frac{V^{3/2}}{x^2},$$

where  $i$  is the current density in amperes per square centimetre, and  $V$  the potential in volts at a distance of  $x$  cm. from the hot plate. It appears that when the current is limited by the space charge it should vary as  $V^{3/2}$  and inversely as  $x^2$ . This agrees approximately with the experimental results. The current is found to increase nearly as  $V^{3/2}$

until it is nearly equal to the saturation current. It then remains nearly constant when  $V$  is increased further.

If instead of two parallel planes, one emitting electrons, we consider two concentric cylinders, the inside one emitting electrons, similar theoretical results are obtained.

Let  $b$  be the radius of the outer cylinder and  $a$  that of the inner one. Let  $V$  be the potential difference between the inner cylinder and a point at a distance  $r$  from its axis. The differential equation which  $V$  must satisfy is then

$$\frac{\partial^2 V}{\partial r^2} + \frac{1}{r} \frac{\partial V}{\partial r} = -4\pi ne.$$

Let now  $i$  denote the current per unit length of the axis of the cylinders, so that

$$i = 2\pi rnev,$$

where  $v$  is the velocity of the electrons,  $n$  the number per unit volume, and  $e$  the charge on one electron. For simplicity we shall suppose that  $-Ve = \frac{1}{2}mv^2$ , which means that we neglect the initial velocity of the electrons. These equations give

$$\frac{\partial^2 V}{\partial r^2} + \frac{1}{r} \frac{\partial V}{\partial r} = -\frac{i}{r} \sqrt{\frac{-2m}{Ve}}.$$

We require a solution of this making  $\partial V/\partial r$  zero at the radius where  $V$  is a minimum, as in the previous problem, and  $V = 0$  at  $r = a$ . When  $a$  is small compared to  $b$ , which is usually the case in practice, then it will be approximately correct to use a solution making  $V = 0$  at  $r = 0$ , provided this solution also makes  $\partial V/\partial r$  small near the inner cylinder. If we assume  $V = Ar^{2/3}$  and substitute this in the differential equation for  $V$ , we find

$$i = -\frac{2}{9} \sqrt{\frac{-2e}{m}} \frac{V^{3/2}}{r}.$$

This satisfies the equation, and makes  $\frac{\partial V}{\partial r}$  small near the inner cylinder as well as  $V = 0$  at  $r = 0$ , and so is an approximate solution of the problem when  $a/b$  is small.

With  $V$  in volts,  $r$  in centimetres, and  $i$  in amperes we get

$$i = -14.65 \times 10^{-6} \frac{V^{3/2}}{b},$$

where  $V$  is now the potential difference between the outer and inner cylinders so that  $r = b$ . This equation is found to agree approximately with the observed currents when  $i$  is less than the saturation current and  $b/a$  is greater than about 10.

It will be observed that when the current is limited by the space

charge it is independent of the temperature of the hot electrode.

Thermionic currents limited by space charge are employed in the three-electrode vacuum tubes now used so extensively as detectors and amplifiers, and for maintaining electrical oscillations.

## 7. Distribution of Electron Velocities.

The distribution of velocities among the electrons emitted by hot metals in a vacuum was investigated by O. W. Richardson and was found to be in agreement with Maxwell's law for the molecules of a gas.

Consider a fine straight wire emitting electrons in a vacuum and surrounded by a concentric cylindrical electrode. Let the number of electrons emitted be small so that space charge effects are negligible. If the potential difference between the wire and the cylinder is  $V$ , then when  $V$  is positive the electrons have to do work  $-Ve$  in order to get across to the cylinder. Let the velocity  $v$  with which an electron is emitted be resolved into two components,  $v_\alpha$  along the axis of the wire, and  $v_r$  along the radius of the cylinder. We suppose the radius of the hot wire very small, so that the electrons can be regarded as starting practically at the axis. The component  $v_\alpha$  does not help the electron to get across and so may be disregarded. For the electron to get across we must have

$$\frac{1}{2}mv_\alpha^2 > -Ve.$$

Let the number of electrons for which  $\frac{1}{2}mv_\alpha^2$  is between  $E$  and  $E + dE$  be given by

$$dn = n f(E) dE,$$

where  $n$  is the number emitted in unit time, so that the saturation current  $i_0 = ne$ .

The current obtained will then be

$$i = i_0 \int_{-V/e}^{\infty} f(E) dE.$$

Differentiating this with respect to  $V$ , we get

$$\frac{di}{dV} = i_0 e f(-Ve).$$

Thus if we observe the current  $i$  obtained with different values of  $V$  we can determine  $di/dV$  and so get values of the function  $f(E)$ .

It was found experimentally that  $\frac{i}{i_0} = e^{-\alpha V}$ , where  $\alpha$  is a constant. The values found for  $\alpha$  were nearly equal to  $-e/kT$ , where  $k$  is the gas constant for one molecule and  $T$  the temperature of the wire. Hence

$$\frac{1}{i_0} \frac{di}{dV} = \frac{e}{kT} e^{Ve/kT} = e f(-Ve),$$

so that

$$f(E) = \frac{1}{kT} e^{-E/kT}.$$

This shows that the electrons have the Maxwell velocity distribution for the velocity component  $v_r$ , for according to Maxwell's law the number of the molecules in unit volume having velocity components  $u$  between  $u$  and  $u + du$  is

$$nA e^{-u^2/2u} du.$$

The number of these going through a unit area perpendicular to  $u$  in unit time is therefore

$$nA e^{-u^2/2u} du,$$

and the total number going through is

$$nA \int_0^{\infty} \epsilon^{-qu^2} u du = \frac{nA}{2q}.$$

Hence the fraction of those going through which have velocities between  $u$  and  $u + du$  is  $2q\epsilon^{-qu^2} u du$ .

But  $q = \frac{m}{2kT}$  (section 5), and  $E = \frac{1}{2}mu^2$ , so that the fraction is equal to  $\frac{1}{kT} \epsilon^{-E/kT} dE$ , in agreement with the experimental result.

Richardson also showed that the velocity components parallel to the emitting surface are distributed in accordance with Maxwell's law.

It appears, therefore, that although the electrons in the metal have very little kinetic energy, yet after escaping they have the same energy as gas molecules at the same temperature. This rather surprising conclusion may perhaps be explained by means of the quantum theory. In the chapter on the quantum theory it is shown that the possible values of the kinetic energy of a monatomic gas molecule contained in a cubical box of volume  $V$  are given by

$$s_n = \frac{h^2}{2m} \left( \frac{3n}{4\pi V} \right)^{2/3},$$

where  $h$  is Planck's constant,  $m$  the mass of one molecule, and  $n = 0, 1, 2, 3$ , &c. Inside the metal the electrons have very short free paths, probably of the order of magnitude of  $10^{-8}$  cm., whereas outside they have long free paths which we may take to be say 1 cm. long. If we suppose, therefore, that  $V$  in the metal is  $10^{-24}$  c. c. and outside 1 c. c., we see that the possible energies outside are  $10^{16}$  times smaller than those inside the metal.

Now  $h = 6.5 \times 10^{-27}$  and  $m = 9 \times 10^{-28}$ , so that

$$s_n = 0.92 \times 10^{-26} \frac{n^{2/3}}{l^2},$$

where  $l$  is the length of the side of the cubical box which we take to correspond with the mean free path. With  $n = 1$  and  $l = 10^{-8}$ , this gives  $s_n = 0.92 \times 10^{-10}$  erg, but, with  $n = 1$  and  $l = 1$ ,  $s_n = 0.92 \times 10^{-26}$  erg. The average energy of a gas molecule at  $T = 1000^\circ$  K. is  $1000 k$ , which is  $1.372 \times 10^{-13}$  erg. Thus in the metal at  $1000^\circ$  K. the smallest possible energy is several hundred times  $kT$ , whereas outside it is very much smaller than  $kT$ . It follows according to the quantum theory that in the metal the free electrons have practically no kinetic energy since nearly all of them must be in the state of zero energy, whereas outside they have the same average energy as ordinary gas molecules.

Under some circumstances hot bodies emit positively charged molecules or ions as well as electrons. This emission of positive ions will be discussed in the chapter on positive rays. It is found that some metallic oxides emit electrons at high temperatures; for example, calcium and barium oxides do this very freely. The emission of electrons by such oxides appears to vary with the temperature in much the same way as in the case of metals.

#### REFERENCE

*The Emission of Electricity from Hot Bodies.* O. W. Richardson.

## CHAPTER IV

# Photo-electricity

### 1. Ultra-violet Light and Emission of Electrons.

Hertz in 1887 discovered that when ultra-violet light falls on a spark gap the sparks pass more easily. This effect was investigated by Hallwachs, Elster and Geitel, and others, and it was found that ultra-violet light causes a negatively charged conductor to lose its charge but has no effect on a positively charged conductor. Lenard made the important discovery that the kinetic energy of the negatively charged particles emitted when ultra-violet light falls on a negatively charged conductor is independent of the intensity of the light, but increases with the frequency of the light waves.

J. J. Thomson and Lenard in 1899 measured the ratio of the charge  $e$  to the mass  $m$  of the charged particles emitted, and found it equal to about  $10^7$  electromagnetic units per gram. This was about the same value of  $e/m$  as had been found for cathode rays or electrons, so that it was clear that ultra-violet light causes solid bodies to emit electrons.

The kinetic energy of the electrons emitted can be determined by finding the potential difference necessary to stop them.

An insulated electrode surrounded by a hollow metal conductor, both contained in a glass bulb which can be exhausted, is illuminated by the ultra-violet light. The electrode emits electrons and so becomes positively charged. The number of electrons which get across from the electrode to the surrounding conductor diminishes as the potential difference increases, and the potential difference soon attains a constant maximum value just sufficient to prevent any more electrons getting across. This maximum potential difference is a measure of the maximum kinetic energy of the electrons emitted. If  $V$  denotes this potential difference and  $e$  the charge on one electron, then  $Ve$  is equal to the maximum kinetic energy. The potential difference can be measured by connecting the electrode and conductor to an electroscope or quadrant electrometer. Experiments of this kind have been made by Ladenburg, A. L. Hughes, Richardson and Compton, and Millikan. Hughes, and Richardson and Compton found that  $Ve$  is of the same order of magnitude as  $h\nu$ ,  $h$  being Planck's constant of the quantum theory and  $\nu$  the frequency of the incident light. Millikan eliminated

various sources of error and obtained much more accurate results than previous workers, so that only his experiments need be considered in detail.

## 2. Millikan's Experiments. Critical Frequency.

In Millikan's experiments a small block of an alkali metal was illuminated by light in a very perfect vacuum. The surface of the block was scraped in the vacuum by a cutter worked by an electromagnet outside. In this way a clean metallic surface was obtained free from any film of oxide. The scraped surface was arranged in front of an insulated cylinder of oxidized copper gauze connected to a quadrant electrometer, and the charge received by this cylinder was observed when the surface was illuminated by monochromatic light of known frequency. The potential difference between the gauze cylinder and illuminated block was increased until it was enough to prevent any electrons from the block reaching the cylinder. The alkali metals emit electrons when exposed to light of much lower frequencies than are necessary to cause copper oxide to emit electrons. Millikan was therefore able to use only light which had no effect on the copper cylinder and so to avoid errors due to the cylinder emitting electrons. The contact potential difference between the alkali metal and copper oxide was determined by moving the metal block in the vacuum to a position opposite a plane copper electrode coated with oxide. This electrode could be moved in and out so as to vary the distance between it and the alkali metal block. The potential difference between this movable electrode and the block was adjusted until moving the electrode in and out made no difference to its potential when it was insulated. The potential difference is then equal to the contact potential difference and there is then no electric field in the space between the block and electrode. The potential difference required to stop the electrons from getting from the block to the gauze cylinder was taken to be the difference between the potential difference observed and the contact potential difference. In this way errors due to the contact potential difference were got rid of.

Millikan found that his results could be represented by the equation  $Ve = h(\nu - \nu_0)$ , where  $V$  is the corrected potential difference required to stop the electrons,  $h$  Planck's constant,  $\nu$  the frequency of the light used, and  $\nu_0$  the smallest frequency which causes any electrons to be emitted. The frequency  $\nu_0$  is called the threshold or critical frequency and it is different for different substances. From his experiments Millikan obtained the value  $h = 6.56 \times 10^{-27}$ , using the value of  $e = 4.774 \times 10^{-10}$  which he had previously determined. This value of  $h$  agrees well with that deduced from measurements of heat radiation and the frequencies of spectral lines, and is believed to be very near the true value.

It is found that the critical frequency for any substance depends

on the state of the surface used. Traces of gases and films of oxide or other impurities greatly affect it. The critical frequency is closely related to the contact potential difference and to the thermionic work function. Consider two parallel plates of different metals, and let them be connected together by a wire. If we disregard the small potential differences of thermoelectricity we may consider the two plates to be at the same potential. There will, nevertheless, be an electric field in the space between the plates equal to the contact potential difference divided by the distance between them. Let  $V_1$  and  $V_2$  be the potentials at the surface of the plates just outside the metals, and  $e\phi_1$  and  $e\phi_2$  the amounts of work required to remove an electron from inside the metals to a point just outside; then we have (see Chap. III, end of section 3).

$$\phi_1 - \phi_2 + V_2 - V_1 = 0.$$

### 3. Einstein's Theory.

According to the theory originally put forward by Einstein, when light falls on a metal electrons in the metal receive quanta  $h\nu$  of energy from the light so that if they escape from the metal their kinetic energy is  $h\nu - e\phi$ , since  $e\phi$  is the work required to remove an electron from the metal,  $\phi$  being the thermionic work function.

Hence  $e\phi = h\nu_0$ , and  $e(V_2 - V_1) = h(\nu_{02} - \nu_{01}) = e(\phi_2 - \phi_1)$ .

These relations appear to be confirmed by measurements of the contact potential differences, critical frequencies, and thermionic work functions, but cannot be regarded as definitely proved to be correct.

It appears that the electrons emitted with the maximum energy  $h(\nu - \nu_0)$  must be the free electrons in the metal, since if electrons were emitted from the interior of atoms we should expect them to come out with energy less than  $h\nu - e\phi$ . Some of the electrons emitted may come from inside atoms, since most of them come out with less than the maximum energy. The light penetrates a short distance into the metal, so that some of the electrons may come out from points inside the metal and may lose energy by collisions before they get out.

Einstein's equation  $\bar{V}e = h(\nu - \nu_0)$  holds good also for the emission of electrons due to X-rays.

The critical frequency  $\nu_0$  for most substances is in the ultra-violet region, but for the alkali metals it is in the visible spectrum or even in the infra-red.

The number of electrons emitted is found to be proportional to the intensity of the light.

### 4. Photo-electricity and the Classical Wave Theory.

It is apparently impossible to explain the phenomena of photo-electricity in a satisfactory manner by means of the classical wave

theory of light. The difficulty is to explain how a very small fraction of the electrons get energy  $h\nu$  and the rest get none. For example, when X-rays are passed through air a few atoms here and there emit electrons with energy  $h\nu$  and the rest are apparently in no way affected.

According to the classical wave theory an electron which has a natural frequency of vibration equal to  $\nu$  can absorb the energy of a train of waves, of frequency  $\nu$ , from an area of the order of that of a square with sides one wave-length long. An ordinary X-ray tube gives out about  $10^8$  ergs of X-ray energy per second, and the ionization due to it can easily be detected at 3000 cm. The energy falling on one square wave length or about  $10^{-15}$  sq. cm. at this distance from the tube is therefore about  $10^{-15}$  ergs per second. The energy with which the electrons are emitted is about  $10^{-7}$  ergs, so that it would take an electron  $10^8$  sec. or 1000 days to absorb it. But when the X-ray tube is started the ionization begins immediately. We might suppose that the energy of the emitted electrons is not derived from the rays but that the rays, so to speak, merely pull a trigger which releases the electron from the atom. However, if this were so we should expect the energy of the electrons emitted to be determined by the nature of the emitting atom, and not by the frequency of the rays as is the case. Moreover, it is difficult to believe that electrons having all possible natural frequencies are present in any substance. It seems certain that the energy of the electrons is derived from the rays, and the classical theory cannot explain how so much of it is concentrated into particular electrons in the time available. According to the quantum theory of radiation the radiation is supposed to consist of quanta, each of energy  $h\nu$ , which travel out from the source with the velocity of light. If an electron absorbs one of these quanta it gets energy  $h\nu$ . This quantum theory, therefore, explains the chief facts of photo-electricity, but of course it is hard to see how it can explain the facts of interference and diffraction which agree so well with the wave theory. This question is also discussed in the chapters on the quantum theory and on X-rays. Here we may say that there is no reason to suppose that a satisfactory explanation of both sets of facts will not eventually be discovered. It is quite unnecessary to adopt a mystical attitude and say that the human mind cannot understand such phenomena. Such an attitude is no satisfactory apology for an unintelligible theory.

### 5. Relation of Emission to Frequency. O. W. Richardson's Theory.

The number of electrons emitted by a metal per unit energy of the incident light depends upon the frequency of the light. It is zero for frequencies less than the critical frequency  $\nu_0$ , and for frequencies greater than  $\nu_0$  it increases at first as  $\nu$  increases, reaches a maximum value, and then decreases. The frequency which gives the maximum emission is about  $\frac{3}{2}\nu_0$  in most cases. With some metals more than one maximum

has been observed. An interesting theory of the variation of the emission with the frequency  $\nu$  of the light has been given by O. W. Richardson. Consider a cylinder and piston maintained at a constant temperature  $T$ . The cylinder will be filled with black body radiation and this will cause the walls to emit electrons. Let the number of electrons per cubic centimetre in the cylinder be  $n$  and let the pressure they exert on the piston be  $p$ . We suppose  $n$  so small that the electric field due to the negative charges on the electrons may be neglected. If we suppose that the thermodynamical theory of evaporation can be applied to the escape of the electrons as in the theory of thermionics we have

$$N = \frac{2\pi m k^2}{h^3} T^2 e^{-w_0/kT},$$

where  $N$  is the number of electrons escaping from unit area in unit time, which is equal to the number absorbed from the electron gas by unit area in unit time. This equation is merely the equation for thermionic emission and the proof of it is given in the chapter on thermionics (Chap. 3, section 5).

Now let the energy density of the radiation between the frequencies  $\nu$  and  $\nu + d\nu$  be denoted by  $E(\nu)d\nu$ . The radiation falling on unit area in unit time is then  $\frac{c}{4} E(\nu)d\nu$ . For the proportion of  $E$  which belongs to rays whose directions lie within a small solid angle  $d\omega$  is  $d\omega/4\pi$ . The radiation falling per unit time on a plane area  $A$  in directions inclined to the normal to  $A$  at angles between  $\theta$  and  $\theta + d\theta$  is therefore  $Ec \cdot A \cos \theta \cdot 2\pi \sin \theta d\theta / 4\pi$ . Integrating from  $\theta = 0$  to  $\theta = \frac{\pi}{2}$ , we get  $\frac{1}{4} cEA$ , the result stated. Let  $F(\nu)$  be the number of electrons liberated from the surface by unit incident light energy of frequencies between  $\nu$  and  $\nu + d\nu$ , so that the number of electrons liberated per unit area per unit time is

$$\frac{c}{4} \int_0^\infty F(\nu) E(\nu) d\nu = N.$$

For  $E(\nu)$  we may use Wien's approximate formula (Chap. V, section 11) since only high frequencies are effective, so that

$$E(\nu) = \frac{8\pi}{c^3} h \nu^3 e^{-h\nu/kT},$$

and we get  $\frac{c}{4} \int_0^\infty F(\nu) \frac{8\pi}{c^3} h \nu^3 e^{-h\nu/kT} d\nu = \frac{2\pi m k^2}{h^3} T^2 e^{-w_0/kT},$

or  $\int_0^\infty F(\nu) \nu^3 e^{-h\nu/kT} d\nu = \frac{c^2 m k^2}{h^4} T^2 e^{-w_0/kT}.$

This equation is satisfied by

$$F(\nu) = \frac{mc^2}{\nu^2 \hbar^2} \left( 1 - \frac{w_0}{\hbar \nu} \right)$$

for  $\nu > w_0/\hbar$ , and  $F(\nu) = 0$  for  $\nu < w_0/\hbar$ . Differentiating  $F(\nu)$  with respect to  $\nu$  and putting  $\frac{dF(\nu)}{d\nu} = 0$ , we find that  $F(\nu)$  has a maximum value when  $\hbar \nu = \frac{3}{2}w_0$ . This agrees with the experimental results that the maximum emission is obtained with a frequency equal to  $\frac{3}{2}$  of the critical frequency, and that there is no emission when  $\nu$  is less than the critical value  $w_0/\hbar$ .

## 6. Thermionic Emission and Photo-electric Action.

In this theory the thermionic emission is not considered separately from the photo-electric emission, so that the theory really involves the assumption that the thermionic emission is caused by photo-electric action of radiation. Measurements of the photo-electric emission due to the radiation from hot tungsten by S. C. Roy (*Proc. Roy. Soc.*, Oct., 1926) show that it is quite possible that thermionic emission may be due entirely to photo-electric action.

The energy of the escaping electrons may be obtained if we assume that the electrons in the cylinder have the same energies as the molecules of a gas at the temperature  $T$ . The thermodynamical theory of evaporation would not be applicable to the electrons if this were not the case.

Let  $T_\nu$  denote the kinetic energy of the electrons liberated by the light of frequency  $\nu$ , so that

$$\frac{c}{4} \int_0^\infty T_\nu F(\nu) E(\nu) d\nu$$

is the total energy of the electrons escaping from unit area in unit time. This may be put equal to the energy of the  $N$  electrons falling on unit area in unit time, which is  $2NkT$  according to the kinetic theory of gases. Using the value found for  $F(\nu)$  we find that

$$T_\nu = \hbar \nu - w_0$$

in agreement with Einstein's equation. This result is not in agreement with the experimental fact that most of the electrons emitted have energy less than  $\hbar \nu - w_0$ , but the discrepancy may be only apparent. We suppose that the electrons receive energy  $\hbar \nu$  from the light initially, so that if they have energy less than  $\hbar \nu - w_0$  they must have lost some energy by collisions or otherwise.

Richardson's theory does not involve the assumption that the light consists of quanta having energy  $\hbar \nu$ , but it is consistent with that view.

When light of sufficiently high frequency is passed through gases electrons are liberated from the gas molecules. The critical frequencies for gases are greater than those for solid bodies. In the case of air the critical wave-length according to A. L. Hughes is about 1350 Å. The ionization of gases by X-rays is due to the emission of electrons by the atoms.

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## CHAPTER V

# The Quantum Theory

### 1. Inadequacy of Newtonian Dynamics.

During the nineteenth century it was generally believed that material phenomena would prove capable of explanation on the old Newtonian system of dynamics. Matter was supposed to consist of minute particles which moved in accordance with Newton's laws of motion. However, certain phenomena, notably those of heat radiation, have emerged which seem to be inconsistent with the laws of classical dynamics, and the quantum theory was put forward by Max Planck as an explanation of such phenomena. This theory has now been applied successfully in several important branches of physics, for example, the theories of spectra, photo-electricity, and chemical equilibrium, and its fundamental character and value are universally recognized. The general validity of Newton's dynamics, therefore, can no longer be admitted. It appears that the laws of motion of bodies consisting of enormous numbers of atoms do not apply to atomic systems.

### 2. Microscopic and Macroscopic States. Statistical Mechanics.

It is necessary to distinguish between macroscopic and microscopic states of a substance. A macroscopic state is one determined by quantities such as pressure and temperature which can be measured by ordinary apparatus. For example, the macroscopic state of a gas is fixed by its pressure and volume. A microscopic state is one determined by the position and motion of all the parts of the substance, however small. Thus, a microscopic state of a gas requires for its specification the positions and velocities of all the molecules of which the gas is believed to be composed.

Quantities such as pressure, internal energy per unit mass, and temperature, which are used to specify the condition of a substance, are average values over very large numbers of atoms. If, for example, we know the temperature and the entropy of a given quantity of any homogeneous substance, then we know that it is in a certain macroscopic state capable of being reproduced and indistinguishable from

other states of the same substance having equal temperatures and specific entropies. But for any such macroscopic state there is an enormous number of different microscopic states. Thus, a gas at a given temperature and pressure is in a definite macroscopic state, but its molecules are moving about so that its microscopic state is continually changing, and in a short time it passes through an enormous number of different microscopic states. The heat radiation inside an empty enclosure the walls of which are maintained at a constant temperature is in a definite macroscopic state. The energy density for any given range of frequencies has a definite value depending only on the temperature of the walls. But to specify the microscopic state of the radiation in the enclosure at any instant it would be necessary to give the strength and direction of the electric and magnetic fields at every point in the enclosure. The radiation in the enclosure is continually moving with the velocity of light, so that its microscopic state is continually changing and it passes through an enormous number of microscopic states in a short time.

It is clear that to any macroscopic state of a substance there corresponds an enormous, perhaps an infinite, number of microscopic states. During the rapid change from one microscopic state to another which continually goes on in material substances which are in a state of equilibrium in a fixed macroscopic state, the macroscopic quantities determining the macroscopic state do not vary perceptibly. The value of such a macroscopic quantity, which can be measured, is an average over a large number of molecules for a short but finite time, or over a large volume, and it remains constant within the limits of error of observation in ordinary cases.

If a solid sphere is immersed in a gas, the uniform pressure of the gas over the surface of the sphere gives no resultant force on the sphere when we consider the action of the gas on the sphere from the macroscopic or large-scale point of view. Microscopically, however, the sphere is not subjected to a uniform pressure but to a series of molecular impacts, and, indeed, if the sphere is very small it does not remain at rest but moves about in an irregular manner, owing to the irregular distribution of the molecular impacts over its surface. This motion, the well-known Brownian movements of small particles immersed in a liquid or gas, is a microscopic motion. Macroscopically, the sphere is to be regarded as at rest in its mean position.

The question arises how it is that the macroscopic state of a substance can remain sensibly constant while its microscopic state is continually changing. The reason must be that of all the possible microscopic states the vast majority correspond to values of the macroscopic quantities differing inappreciably from the constant values. If any microscopic state have an appreciably different value of the macroscopic quantities, the chance of its lasting long enough for the

change in a macroscopic quantity to be observable is therefore negligible.

When a substance is not in a state of macroscopic equilibrium its state changes until equilibrium is reached. The relation of such changes to the microscopic states will now be considered. Since the number of microscopic states corresponding to any state differing appreciably from the state of macroscopic equilibrium must be very small compared with the whole number of possible microscopic states, the chance of the substance remaining an appreciable time in such a state must also be very small, and the substance changes to the equilibrium macroscopic state because this state corresponds to nearly all possible microscopic states. The change from a state which is not one of equilibrium to an equilibrium state thus involves an increase in the number of possible microscopic states. It is clear, therefore, that an equilibrium state is one for which the number of microscopic states is a maximum. Such considerations, of course, only apply to substances or systems the microscopic parts of which are continually changing from one microscopic state to another, so that in the course of time the system may be supposed to pass through all possible microscopic states.

It appears then that when a substance consists of a very great number of individuals, the behaviour of the substance as a whole must be investigated by methods of statistics.

### 3. Entropy and Probability.

Let  $W$  denote the number of possible microscopic states through which the substance may pass in the course of time while in a given macroscopic state. Then the condition of equilibrium is  $W = \text{maximum}$  or  $\delta W = 0$ . If the microscopic parts of the system are at rest so that its microscopic state does not change with time then  $W = 1$ . According to the second law of thermodynamics the entropy  $\Phi$  of any isolated system is increased by any spontaneous change which takes place in the system, and in a state of equilibrium the entropy is a maximum. Thus it appears that both  $W$  and  $\Phi$  tend to increase to maximum values.  $W$  may be called the thermodynamical probability of the state of the system, and we may then say that the system changes to a more probable state unless it is in a state of maximum probability. The second law of thermodynamics evidently expresses the same thing in terms of  $\Phi$  instead of  $W$ , so that we should expect  $\Phi$  to depend on  $W$ . This idea was first put forward by Boltzmann. Let us then suppose that the entropy  $\Phi$  is some function of  $W$ , or let

$$\Phi = f(W).$$

Now consider two entirely separate systems having entropies  $\Phi_1$  and  $\Phi_2$  and thermodynamic probabilities  $W_1$  and  $W_2$ . Then

$$\Phi_1 = f(W_1), \quad \Phi_2 = f(W_2).$$

The total entropy of the two systems is  $\Phi_1 + \Phi_2$ , and since for each microscopic state of the first system there are  $W_2$  states of the second, the thermodynamic probability of the two systems considered together as one must be the product  $W_1 W_2$ . Thus we have

$$\Phi_1 + \Phi_2 = f(W_1 W_2)$$

so that

$$f(W_1) + f(W_2) = f(W_1 W_2).$$

Differentiating this equation with respect to  $W_1$ , and also with respect to  $W_2$ , we obtain

$$W_1 f'(W_1) = W_2 f'(W_2) = W_1 W_2 f'(W_1 W_2).$$

We may suppose  $W_1$  to change while  $W_2$  remains constant, since the two systems were supposed entirely separate, so that we must have

$$W_1 f'(W_1) = W_2 f'(W_2) = \text{constant}.$$

Hence  $f(W) = k \log W + C$  where  $k$  is a universal constant, the same for all systems, and  $C$  is another constant, not necessarily the same for all systems. Hence  $\Phi = k \log W + C$  for any system. In classical thermodynamics the entropy is taken to be the difference between the entropy in the actual state and that in an arbitrarily chosen standard state. If  $W_0$  is the thermodynamical probability in the standard state then  $\Phi = k \log W - k \log W_0 = k \log (W/W_0)$ .

The number of possible microscopic states  $W$  of a system corresponding to a given macroscopic state might be expected to be infinite. Thus if the velocity of an atom can vary continuously it has an infinite number of possible values, so that a gas at a given pressure and temperature should have an infinite number of possible microscopic states. But if  $W$  is infinite then  $\Phi = k \log W + C$  must also be infinite. We might suppose that though  $W$  and  $W_0$  were both infinite their ratio  $W/W_0$  could have a definite finite value, but this method of avoiding the difficulty has not proved satisfactory. Planck supposes that  $W$  is not infinite but has a definite value for any system, and also that the constant  $C$  is equal to zero so that  $\Phi = k \log W$ . This leads to an absolute value of the entropy and to the quantum theory. It follows that a system in a given macroscopic state can only exist in a finite number of microscopic states. Thus, for example, the velocities of the molecules of a gas cannot vary continuously, but must be restricted in some way to a finite number of possible values, or at least changes in the velocities less than certain finite amounts must be supposed not to constitute a change in the microscopic state.

#### 4. State Space of a System.

Consider a system of any kind, for example, an atom of any element. Let the microscopic state of the system be determined by co-ordinates  $q_1, q_2, \dots, q_n$  and the corresponding momenta  $p_1, p_2, \dots, p_n$ ; that is, let  $p_m$  be the momentum associated with the co-ordinate  $q_m$ . The  $2n$  quantities  $p_1, p_2, \dots, p_n$  and  $q_1, q_2, \dots, q_n$  may be regarded as the co-ordinates of a point in a space of  $2n$  dimensions. This representative point will move along a path which in the case of a periodic motion of the system will be a closed curve. So long as the energy of the system remains unchanged the point will continue to move round and round this closed curve, and if we regard the state of the system as determined by its energy then the motion of the representative point round the curve does not involve a change in the state of the system. The space of  $2n$  dimensions is called the state space of the system.

Suppose now that we have a large number  $\mathcal{N}$  of such systems all alike so that the state of each one can be represented in the space of  $2n$  dimensions by a representative point. The  $\mathcal{N}$  points will all describe curves in the space. The microscopic state of the collection may be regarded as determined by the distribution of energy among the  $\mathcal{N}$  systems. In the state space we can imagine surfaces or regions drawn in which the energy is constant. Then all the systems having the same energy will have representative points moving on the same surface of constant energy. According to classical dynamics we should expect that the energy of the system could vary continuously, so that there would be an infinite number of regions or surfaces of constant energy on which the representative points could move. The number of possible microscopic states of the collection of  $\mathcal{N}$  systems would therefore be infinite. In the quantum theory it is supposed that the number of possible microscopic states is finite, so that only certain definite values of the energy of a system are possible. The surfaces of constant energy corresponding to these possible values will divide the state space up into finite regions, and all the representative points will be on these surfaces.

When the energy of one of the systems changes from one possible value to another, its representative point is supposed to jump from one of the surfaces to another without occupying the intermediate positions effectively. Such jumps are supposed to be due to actions between the systems or between the systems and the radiation in the collection of  $\mathcal{N}$  systems. It is important to remember that any collection of atoms always contains a certain amount of energy in the form of radiation. This radiant energy is generally small compared with the energy of the atoms and so can be neglected in many cases, but it provides a reservoir of energy into which or from which the atoms can give out or absorb energy.

### 5. Planck's Theory of Entropy and Free Energy.

Let  $\epsilon_1, \epsilon_2, \epsilon_3 \dots$  denote the possible values of the energy of a system, so that the total energy of the  $\mathcal{N}$  similar systems or parts of the collection is given by

$$E = N_1\epsilon_1 + N_2\epsilon_2 + \dots = \Sigma N\epsilon,$$

where  $N_1, N_2 \dots$  denote the numbers of the parts having energies  $\epsilon_1, \epsilon_2 \dots$  respectively.

If two systems having energies  $\epsilon_1$  and  $\epsilon_2$  act on each other so that their energies change to  $\epsilon_1'$  and  $\epsilon_2'$ , then if  $\epsilon_1 + \epsilon_2$  is not equal to  $\epsilon_1' + \epsilon_2'$  the difference may be supposed supplied from the radiation present. Also if the energy of a system changes spontaneously from  $\epsilon_1$  to  $\epsilon_1'$  the difference  $\epsilon_1 - \epsilon_1'$  may be supposed emitted in the form of radiation. The total radiant energy is small compared with  $E$ , and its average value remains constant so long as  $E$  is not changed, but it must fluctuate about its average value owing to the emission and absorption as the systems change from one possible value of  $\epsilon$  to another. The number  $W$  of possible microscopic states corresponding to a given distribution of the energy is given by

$$W = \frac{\mathcal{N}!}{N_1! N_2! N_3! \dots}.$$

For if all the  $\mathcal{N}$  parts got different amounts of energy the number of possible arrangements would be equal to the number of permutations of  $\mathcal{N}$  different objects, each permutation containing all the objects, or  $\mathcal{N}!$ . When  $N_1$  of the objects are identical, let the number of different permutations be  $x$ . By changing the  $N_1$  identical objects to  $N_1$  different objects,  $N_1!$  different permutations could be made out of each of the  $x$  permutations, so that

$$xN_1! = \mathcal{N}!.$$

In the same way it is easy to see that when the objects are divided into groups containing  $N_1, N_2, \&c.$ , identical objects the number of permutations is equal to the expression given above for  $W$ .

An approximate value of  $N!$ , sufficiently exact for the purpose of calculating  $\log W$  when  $\mathcal{N}, N_1, N_2, \&c.$ , are large, may be easily obtained. We have

$$\log N! = \log 1 + \log 2 + \log 3 + \log 4 + \dots + \log N,$$

$$\text{and} \quad \log N^N = \log N + \log N + \dots + \log N,$$

$$\text{so that} \quad \log(N!/N^N) = \log \frac{1}{N} + \log \frac{2}{N} + \dots + \log \frac{N}{N}.$$

Put  $\frac{1}{N} = dx$ , so that

$$\begin{aligned}\frac{1}{N} \log(N!/N^N) &= \log dx \cdot dx + \log 2dx \cdot dx + \dots + \log Ndx \cdot dx \\ &= \int_0^1 \log x \cdot dx = -1.\end{aligned}$$

Hence  $\log N! = N \log N - N = \log(N/e)^N$ , so that  $N! = \left(\frac{N}{e}\right)^N$ . Using this value for  $N!$ , we get

$$W = \frac{(\mathcal{N}/e)^{\mathcal{N}}}{(N_1/e)^{N_1}(N_2/e)^{N_2}\dots} = \frac{\mathcal{N}^{\mathcal{N}}}{N_1^{N_1}N_2^{N_2}N_3^{N_3}\dots},$$

since  $N_1 + N_2 + \dots = \mathcal{N}$ .

Hence  $\log W = \mathcal{N} \log \mathcal{N} - \sum N \log N$ .

Now let  $w_1 = N_1/\mathcal{N}$ ,  $w_2 = N_2/\mathcal{N}$ , &c., so that  $\sum w = 1$ , and  $\log W = \mathcal{N} \log \mathcal{N} - \sum \mathcal{N} w \log \mathcal{N} w$ ,

or  $\log W = -\mathcal{N} \sum w \log w$ .

The entropy  $\Phi$  of the system is then given by

$$\Phi = k \log W = -k \mathcal{N} \sum w \log w.$$

To find the distribution of the energy in the system in the state of equilibrium, we make  $\Phi$  a maximum subject to the conditions  $\sum w = 1$  and  $E = \mathcal{N} \sum w \epsilon = \text{constant}$ .

Thus  $\delta \Phi = 0 = \sum \delta w \log w + \sum \delta w = \sum \delta w \log w$ ,  
 $\delta E = 0 = \sum \epsilon \delta w$ , and  $\sum \delta w = 0$ .

Hence  $\sum (\log w + \beta \epsilon + \gamma) \delta w = 0$ ,

where  $\beta$  and  $\gamma$  are undetermined multipliers. If  $\beta$  and  $\gamma$  are properly chosen, this equation will be true for any values of the  $\delta w$ 's, so that we have

$$\log w_n = -\beta \epsilon_n - \gamma,$$

or  $w_n = \alpha e^{-\beta \epsilon_n}$ . . . . . (1)

Here  $\alpha$  and  $\beta$  are constants having the same values for all parts of the system. The entropy in the equilibrium state is therefore given by

$$\Phi = k \mathcal{N} \sum w (\beta \epsilon - \log \alpha),$$

or  $\Phi = k \mathcal{N} \beta \sum w \epsilon - k \mathcal{N} \log \alpha$ ,

so that  $\Phi = k \beta E + k \mathcal{N} \log \sum e^{-\beta \epsilon_n}$ , . . . . . (2)

since  $\sum w_n = \alpha \sum e^{-\beta \epsilon_n} = 1$ , and  $E = \mathcal{N} \sum w_n \epsilon_n$ .

The thermodynamical definition of  $\Phi$  gives

$$\delta\Phi = \frac{\delta E + p\delta V}{\theta},$$

where  $p$ ,  $V$ , and  $\theta$  denote the pressure, volume, and absolute temperature of the system. But, regarding  $\Phi$  as a function of  $E$  and  $V$ , we have

$$\delta\Phi = \left(\frac{\partial\Phi}{\partial E}\right)_V \delta E + \left(\frac{\partial\Phi}{\partial V}\right)_E \delta V,$$

so that

$$\left(\frac{\partial\Phi}{\partial E}\right)_V = \frac{1}{\theta} \text{ and } \left(\frac{\partial\Phi}{\partial V}\right)_E = \frac{p}{\theta}.$$

Differentiating the equation  $\Phi = k\beta E + k\mathcal{N} \log \sum e^{-\beta\epsilon_n}$  with respect to  $\beta$ , we get

$$\frac{\partial\Phi}{\partial\beta} = kE + k\beta \frac{\partial E}{\partial\beta} - \frac{k\mathcal{N} \sum \epsilon_n e^{-\beta\epsilon_n}}{\sum e^{-\beta\epsilon_n}}.$$

But 
$$\frac{\mathcal{N} \sum \epsilon_n e^{-\beta\epsilon_n}}{\sum e^{-\beta\epsilon_n}} = \mathcal{N} \alpha \sum \epsilon_n e^{-\beta\epsilon_n} = \mathcal{N} \sum w_n \epsilon_n = E,$$

so that 
$$\frac{\partial\Phi}{\partial\beta} = k\beta \frac{\partial E}{\partial\beta}.$$

Hence 
$$\frac{\partial\Phi}{\partial E} = \frac{\partial\Phi}{\partial\beta} \frac{\partial\beta}{\partial E} = k\beta = \frac{1}{\theta}.$$

This enables us to introduce the temperatures into the expression (2) for the entropy, and obtain

$$\Phi = \frac{E}{\theta} + k\mathcal{N} \log \sum e^{-\epsilon_n/k\theta}. \quad \dots \quad (3)$$

The free energy  $F$  of the system is defined to be  $E - \theta\Phi$ , so that

$$F = -k\mathcal{N} \theta \log \sum e^{-\epsilon_n/k\theta}.$$

This gives 
$$\theta \frac{\partial F}{\partial \theta} = -k\mathcal{N} \theta \log \sum e^{-\epsilon_n/k\theta} - \frac{k\mathcal{N} \theta^2}{k\theta^2} \frac{\sum \epsilon_n e^{-\epsilon_n/k\theta}}{\sum e^{-\epsilon_n/k\theta}},$$

or 
$$E = F - \theta \frac{\partial F}{\partial \theta}. \quad \dots \quad (4)$$

This is the well-known Gibbs-Helmholtz equation, from which the thermodynamical theory of the system may be deduced. Thus we see that the quantum theory is consistent with classical thermodynamics.

Since  $E = F + \theta\Phi$ , equation (4) gives 
$$\Phi = -\frac{\partial F}{\partial \theta}.$$

## 6. Planck's Constant.

In order to use (3) to calculate the entropy of a system it is of course necessary to determine the possible values  $\epsilon_n$  of the energies of the  $\mathcal{N}$  independent parts. This may require some knowledge of the physical properties of the parts and involves further assumptions characteristic of the quantum theory. The surfaces of constant energy for the possible values of the energy divide the state space into certain finite regions. In the case of periodic systems determined by only one co-ordinate  $q$  and the corresponding momentum  $p$ , the state space is a plane and the closed curves of possible constant energies are supposed by Planck to divide the plane into equal areas  $h$ . The area of the closed curve of energy  $\epsilon_n$  is then equal to  $n h$ , so that

$$\int dp dq = \int p dq = n h,$$

where  $n = 0, 1, 2, 3, \dots$

The quantity  $h$  is called *Planck's constant*. It appears to have the same value in all such cases, and its value has been determined by comparing results deduced from the quantum theory with experimental values. The most probable value of  $h$  is  $6.55 \times 10^{-27}$  erg-seconds. The dimensions of  $h$  are (energy)  $\times$  (time), or action, for

$$(m\dot{x})dx = (m\dot{x}^2)dt.$$

## 7. Monatomic Gas.

As an example, consider the case of  $\mathcal{N}$  atoms of a monatomic gas, each of mass  $m$ , and each contained in a separate cubical box of volume  $V$ . Let the co-ordinates of an atom measured along the edges of its box be  $x, y, z$ , and the corresponding momenta  $m\dot{x}, m\dot{y}$ , and  $m\dot{z}$ . The state space will be of six dimensions, and the representative points of all the atoms having kinetic energies equal to  $s$  will lie on the surface of a sphere of radius  $r$  given by

$$\begin{aligned} r^2 &= (m\dot{x})^2 + (m\dot{y})^2 + (m\dot{z})^2, \\ s &= \frac{1}{2}m(\dot{x}^2 + \dot{y}^2 + \dot{z}^2), \end{aligned}$$

so that

$$r = \sqrt{2sm},$$

and they all also lie inside the volume  $V$ . The six-dimensional volume enclosed by the surface of constant kinetic energy  $s$  is therefore equal to

$$\frac{4}{3}\pi(2sm)^{3/2}V.$$

We shall suppose that the surfaces of constant energy corresponding to possible values of  $s$  divide the state space into equal regions  $h^3$ , since in this case each system has three co-ordinates. Hence

$$\frac{4}{3}\pi(2s_n m)^{3/2}V = n h^3,$$

where  $n = 0, 1, 2, 3, \dots$

The total energy of an atom  $\epsilon_n$  may be taken equal to  $s_n + \epsilon_0$ , where  $\epsilon_0$  denotes its internal energy. The free energy of the collection of  $\mathcal{N}$  atoms in the equilibrium state of maximum entropy can now be calculated by means of the equation

$$F = -k\mathcal{N}\theta \log \Sigma e^{-\epsilon_n/k\theta}.$$

The values of  $s_n$  are very small unless  $n$  is very large, so that unless the total energy  $E$  of the  $\mathcal{N}$  atoms is very small we may replace  $\sum e^{-\epsilon_n/k\theta} = e^{-\epsilon_0/k\theta} \sum e^{-s_n/k\theta}$  by an integral.

We have 
$$s_n = \frac{h^2}{2m} \left( \frac{3n}{4\pi V} \right)^{2/3} = \alpha k\theta n^{2/3},$$

where  $\alpha$  is a constant. Then

$$\sum e^{-s_n/k\theta} = \sum e^{-\alpha n^{2/3}} = \int_0^\infty e^{-\alpha n^{2/3}} dn.$$

Put  $x = n^{1/3}$  so that  $dx = \frac{1}{3}n^{-2/3}dn$ , and

$$\int_0^\infty e^{-\alpha n^{2/3}} dn = 3 \int_0^\infty x^2 e^{-\alpha x^2} dx = \frac{3}{4} \sqrt{\frac{\pi}{\alpha^3}}.$$

Hence 
$$\sum e^{-\epsilon_n/k\theta} = \frac{(2\pi mk\theta)^{3/2} V e^{-\epsilon_0/k\theta}}{h^3}.$$

The free energy of the  $\mathcal{N}$  atoms in separate boxes is therefore given by

$$F = -k\mathcal{N}\theta \log \{ (2\pi mk\theta)^{3/2} V/h^3 \} + \mathcal{N}\epsilon_0,$$

and the entropy  $\Phi = -\partial F/\partial\theta$  by

$$\Phi = k\mathcal{N} \log \{ (2\pi mk\theta)^{3/2} V/h^3 \} + \frac{3}{2}k\mathcal{N}. \quad \dots (5)$$

The energy  $E$  of the  $\mathcal{N}$  atoms is then given by

$$E = F - \theta \frac{\partial F}{\partial\theta} = \mathcal{N}(\frac{3}{2}k\theta + \epsilon_0),$$

so that it appears that  $\frac{3}{2}k\theta$  is the average kinetic energy of one atom. If we now suppose the  $\mathcal{N}$  atoms all contained in one box of volume  $V$ , then if the  $\mathcal{N}$  atoms are all different the number of ways in which the energy elements can be arranged among the atoms will be the same as before, so that the entropy will still be given by equation (5).

If the  $\mathcal{N}$  atoms are all different, the number of ways in which they can be arranged among  $\mathcal{N}$  quantities of energy is  $\mathcal{N}!$ , so that when they are all alike and it makes no difference which particular atom has a given amount of energy the number of possible arrangements is diminished  $\mathcal{N}!$  times, and  $k \log \mathcal{N}!$ , or  $k\mathcal{N} \log \left( \frac{\mathcal{N}}{e} \right)$ , must be subtracted from the entropy given by (5). Hence we get for the entropy of a monatomic gas

$$\Phi = k\mathcal{N} \log \left\{ \frac{V e^{5/2}}{\mathcal{N} h^3} (2\pi mk\theta)^{3/2} \right\}.$$

Differentiating this with respect to  $V$ , keeping  $E$  and therefore  $\theta$  constant, we get

$$\left( \frac{\partial \Phi}{\partial V} \right)_E = \frac{p}{\theta} = \frac{k\mathcal{N}}{V},$$

so that  $pV = k\mathcal{N}\theta$ .

This equation expresses the well-known relation between the pressure, volume, and temperature of a gas, and shows that  $k$  is equal to the gas constant for one molecule. The energy of the gas is the same as that of the  $\mathcal{N}$  atoms in separate boxes, so that

$$E = \frac{3}{2}k\mathcal{N}\theta + \mathcal{N}\epsilon_0,$$

and the free energy is given by

$$F = E - \theta \Phi = -k\mathcal{N}\theta \log \left\{ \frac{V_e}{\mathcal{N} \hbar^3} (2\pi mk\theta)^{3/2} \right\} + \mathcal{N}\varepsilon_0.$$

The distribution of the energy among the atoms of gas is given by the equation

$$w_n = \alpha e^{-\varepsilon_n/k\theta},$$

which when  $\varepsilon_{n+1} - \varepsilon_n$  is very small expresses Maxwell's law. Thus we have obtained the principal results of the kinetic theory of gases by means of the quantum theory.

## 8. Vapour Pressure.

The latent heat of evaporation,  $L$ , of any solid or liquid at a constant temperature  $\theta$  is given by the equation

$$\frac{L}{\theta} = \Phi_1 - \Phi_2,$$

where  $\Phi_1$  denotes the entropy of the vapour and  $\Phi_2$  that of the solid or liquid. At low temperatures  $\Phi_2$  is small and the vapour may be regarded as a perfect gas, so that for substances giving monatomic vapours we have

$$\frac{L}{\theta} = k\mathcal{N} \log \left\{ \frac{V_e^{5/2}}{\mathcal{N} \hbar^3} (2\pi mk\theta)^{3/2} \right\},$$

where  $L$  is the heat of evaporation of  $\mathcal{N}$  atoms. Putting  $V = k\mathcal{N}\theta/p$  and solving for  $\log p$  we get

$$\log p = -\frac{L}{k\mathcal{N}\theta} + \log \left\{ \left( \frac{2\pi m}{\hbar^2} \right)^{3/2} (k\theta)^{5/2} \right\}.$$

This equation has been found to agree very well with the observed vapour pressures of mercury, argon, helium, and other monatomic substances at low temperatures when the values of  $\hbar$  and  $k$  derived from experiments on heat radiation are substituted in it. It may be written

$$\log p = -\frac{L}{k\mathcal{N}\theta} + \frac{5}{2} \log \theta + \frac{5}{2} + i, \quad . . . . . (6)$$

where

$$i = \log \left\{ \left( \frac{2\pi}{\hbar^2} \right)^{3/2} k^{5/2} \right\} + \frac{5}{2} \log m$$

is Nernst's "chemical constant" for a monatomic substance. The pressure of electron gas in equilibrium with a hot metal may also be calculated by (6) if the heat energy absorbed when  $\mathcal{N}$  electrons escape from the metal is substituted for  $L$ , and for  $m$  the mass of one electron. This important application of the quantum theory is discussed in the chapter on thermionics (Chap. III, section 5).

## 9. Simple Oscillators.

We will now consider the case of a system of a large number  $\mathcal{N}$  of oscillators, each consisting of a particle of mass  $m$ , which can only move along a straight line under the action of a force proportional to its distance from a fixed point in the line and directed toward the fixed point. If  $x$  denotes the distance of the particle from the fixed point, then  $m\ddot{x} = -\mu x$ , where  $\mu$  is a constant. A solution of this equation is

$x = A \sin 2\pi\nu t$ , where  $A$  is the amplitude of the oscillations of the particle and  $\nu$  the frequency or number of vibrations in unit time. Substituting this value of  $x$  in the equation  $m\ddot{x} = -\mu x$  we find

$$\nu = \frac{1}{2\pi} \sqrt{\frac{\mu}{m}}.$$

The energy  $\epsilon$  of the vibrating particle is given by

$$\epsilon = \frac{1}{2} m \dot{x}^2 + \frac{1}{2} \mu x^2 + \epsilon_0,$$

where  $\epsilon_0$  stands for any energy it may have when at rest in its equilibrium position. Putting  $m\dot{x} = y$ , we get

$$1 = \frac{y^2}{2m(\epsilon - \epsilon_0)} + \frac{\mu x^2}{2(\epsilon - \epsilon_0)}.$$

This relation between  $x$  and  $y$  is represented by an ellipse in the  $xy$  plane. The representative point of the oscillator moves round the ellipse, making  $\nu$  revolutions in unit time. The area of the ellipse is

$$2\pi(\epsilon - \epsilon_0) \sqrt{\frac{m}{\mu}},$$

which is equal to  $(\epsilon - \epsilon_0)/\nu$ .

According to the quantum theory the possible paths of the particle in the  $xy$  plane will divide the plane into parts of equal area  $h$ . The possible paths will be a series of ellipses having areas  $0, h, 2h, 3h$ , &c., so that the possible values of the energy are given by  $\epsilon - \epsilon_0 = n h \nu$ , where  $n = 0, 1, 2, 3$ , &c. The free energy of the system of  $\mathcal{N}$  oscillators when the energy is distributed so that the entropy has the maximum value is given (section 5) by  $F = -k\mathcal{N}\theta \log \sum e^{-\epsilon_n/k\theta}$ .

$$\begin{aligned} \text{We have } \sum e^{-\epsilon_n/k\theta} &= e^{-\epsilon_0/k\theta} \{1 + e^{-h\nu/k\theta} + e^{-2h\nu/k\theta} + \dots\} \\ &= \frac{e^{-\epsilon_0/k\theta}}{1 - e^{-h\nu/k\theta}}, \end{aligned}$$

$$\text{so that } F = k\mathcal{N}\theta \log (1 - e^{-h\nu/k\theta}) + \mathcal{N}\epsilon_0.$$

Hence the total energy  $E$  of the  $\mathcal{N}$  oscillators in the state of maximum entropy is equal to

$$E = F - \theta \frac{\partial F}{\partial \theta} = \frac{\mathcal{N}h\nu}{e^{h\nu/k\theta} - 1} + \mathcal{N}\epsilon_0.$$

The average energy of vibration,  $\epsilon - \epsilon_0$ , of one oscillator is therefore  $h\nu/(e^{h\nu/k\theta} - 1)$ , and the fraction  $w_n$  of the  $\mathcal{N}$  oscillators which have energy  $\epsilon_n - \epsilon_0 = n h \nu$  is equal to  $e^{-\epsilon_n/k\theta} / \sum e^{-\epsilon_n/k\theta}$ , or  $e^{-n h \nu/k\theta} (1 - e^{-h\nu/k\theta})$ . The fraction which has no energy is  $1 - e^{-h\nu/k\theta}$ . At high temperatures when  $h\nu/k\theta$  is small the average vibrational energy per oscillator is

$k\theta$ , since  $e^{h\nu/k\theta} = 1 + h\nu/k\theta$  when  $h\nu/k\theta$  is small, and the fraction having zero vibrational energy is  $h\nu/k\theta$ , which is small. At low temperatures when  $h\nu/k\theta$  is large the average energy is much less than  $h\nu$ , and nearly all the oscillators have zero vibrational energy. The heat capacity of the system of oscillators is given by

$$\frac{dE}{d\theta} = \frac{\mathcal{N} (h\nu)^2 e^{h\nu/k\theta}}{k\theta^2 (e^{h\nu/k\theta} - 1)^2}.$$

This makes the heat capacity very small at low temperatures and equal to  $\mathcal{N}k$  at high temperatures.

It appears that the average energy per oscillator in the equilibrium state of maximum entropy depends only on the temperature and the frequency of the oscillators. It is easy to see that this result will apply to any collection of oscillators, even if they are not all similar, provided the energy of each is determined by one co-ordinate and the corresponding momentum. Thus if we consider a solid body and suppose that it possesses  $\mathcal{N}_1$  modes of vibration of frequency  $\nu_1$ ,  $\mathcal{N}_2$  of frequency  $\nu_2$ , &c., the energy of its vibrations will be equal to

$$\frac{\mathcal{N}_1 h\nu_1}{e^{h\nu_1/k\theta} - 1} + \frac{\mathcal{N}_2 h\nu_2}{e^{h\nu_2/k\theta} - 1} + \dots = \Sigma \frac{\mathcal{N}_n h\nu_n}{e^{h\nu_n/k\theta} - 1}.$$

## 10. Quantum Theory of Specific Heat.

In this way Einstein and Debye have worked out a quantum theory of the specific heats of solid bodies. We may regard the heat energy of the solid as the energy of elastic waves travelling through it like sound waves through air. The possible frequencies of vibration are the frequencies of stationary waves possible in the solid.

Consider a cube of the solid with sides of length  $a$ , and let  $AB$  (fig. 1) be a plane wave travelling in the direction  $CN$ . Draw a plane  $A'B'$  at a distance from  $AB$  of one-half wave-length. Take the origin  $O$  at one corner of the cube and axes  $x, y, z$  along its edges. Let the cosines of the angles between the direction of propagation of the plane wave and the axes be  $l, m, n$ . Let  $EF$  be a line parallel to the  $y$  axis cutting the planes  $AB$  and  $A'B'$  at  $C$  and  $C'$ . Then

$$CC' = \frac{\lambda}{2m}.$$

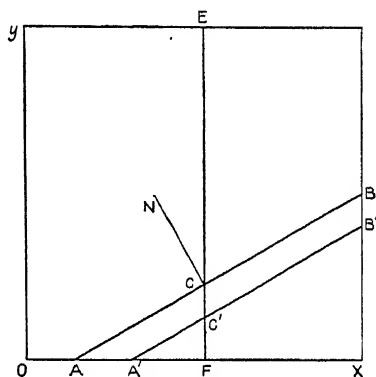


Fig. 1

If the wave  $AB$  is to be one of the systems of standing waves of length  $\lambda$  we must have  $n_2 CC' = a$ , where  $n_2$  is a positive integer. Hence

$$\frac{n_2 \lambda}{2m} = a.*$$

In the same way, considering lines like  $EF$  parallel to the  $x$  and  $z$  axes

we get 
$$\frac{n_1 \lambda}{2l} = a, \text{ and } \frac{n_3 \lambda}{2n} = a.$$

Hence 
$$l^2 + m^2 + n^2 = 1 = \left(\frac{\lambda}{2a}\right)^2 (n_1^2 + n_2^2 + n_3^2).$$

If  $v$  is the velocity of propagation of the waves and  $\nu$  the frequency, then  $\nu \lambda = v$ , so that

$$n_1^2 + n_2^2 + n_3^2 = \left(\frac{2a}{v} \nu\right)^2.$$

Now let  $n_1, n_2, n_3$  be the rectangular co-ordinates of points, so that there will be one point in each unit volume. The number of possible frequencies between 0 and  $\nu$  is thus equal to one-eighth of the volume of a sphere of radius  $\frac{2a\nu}{v}$ , or  $\frac{1}{8} \cdot \frac{4}{3} \pi \left(\frac{2a\nu}{v}\right)^3$ , and the number of possible frequencies between 0 and  $\nu$  per unit volume is equal to  $\frac{4}{3} \pi \left(\frac{\nu}{v}\right)^3$ .

In a solid there will be longitudinal waves and also transverse waves, and the latter can be regarded as forming two sets polarized in perpendicular planes. The total number of frequencies between  $\nu$  and  $\nu + d\nu$  in an elastic solid is therefore

$$4\pi\nu^2 \left(\frac{2}{v_1^3} + \frac{1}{v_2^3}\right) d\nu,$$

where  $v_1$  is the velocity of transverse, and  $v_2$  that of longitudinal waves.

The energy of the vibrations in unit volume between the frequencies 0 and  $\nu$  is therefore

$$4\pi\hbar \left(\frac{2}{v_1^3} + \frac{1}{v_2^3}\right) \int_0^\nu \frac{\nu^3 d\nu}{e^{h\nu/k\theta} - 1}.$$

If the solid contains  $\mathcal{N}$  atoms in unit volume their positions could be determined by  $3\mathcal{N}$  co-ordinates, so that we should not expect more than  $3\mathcal{N}$  possible modes of vibration per unit volume in the solid. Debye therefore supposes that the maximum possible frequency  $\nu_m$  is given by

$$3\mathcal{N} = 4\pi \left(\frac{2}{v_1^3} + \frac{1}{v_2^3}\right) \int_0^{\nu_m} \nu^2 d\nu = \frac{4}{3} \pi \left(\frac{2}{v_1^3} + \frac{1}{v_2^3}\right) \nu_m^3.$$

\* Cf. J. K. Roberts, *Heat and Thermodynamics*, Chap. XX, section 2.

The total energy  $E$  of the vibrations in unit volume is then given by

$$E = 4\pi h \left( \frac{2}{v_1^3} + \frac{1}{v_2^3} \right) \int_0^{\nu_m} \frac{\nu^3 d\nu}{e^{h\nu/k\theta} - 1}.$$

If we put  $h\nu/k\theta = x$ , and  $\psi = h\nu_m/k$ , this equation for  $E$  becomes

$$E = 9\mathcal{N}k \frac{\theta^4}{\psi^3} \int_0^{\psi/\theta} \frac{x^3 dx}{e^x - 1}.$$

The heat capacity of  $\mathcal{N}$  atoms is given by

$$\frac{\partial E}{\partial \theta} = 9\mathcal{N}k \left\{ 4 \left( \frac{\theta}{\psi} \right)^3 \int_0^{\psi/\theta} \frac{x^3 dx}{e^x - 1} - \frac{\psi}{\theta} \frac{1}{e^{\psi/\theta} - 1} \right\}. \quad (7)$$

When  $\theta$  is very large this reduces to  $\frac{\partial E}{\partial \theta} = 3\mathcal{N}k$ , which agrees with Dulong and Petit's law. At low temperatures  $\psi/\theta$  becomes large, so that

$$\frac{\partial E}{\partial \theta} = 36\mathcal{N}k \left( \frac{\theta}{\psi} \right)^3 \int_0^\infty \frac{x^3 dx}{e^x - 1} = \frac{15}{8} \pi^4 \mathcal{N}k \left( \frac{\theta}{\psi} \right)^3. \quad (8)$$

Thus at low temperatures the atomic heat is proportional to the cube of the absolute temperature. Equation (7) shows that the atomic heat capacity is the same function of  $\theta/\psi$  for all solid substances.

$\psi$  is called the *characteristic temperature* of the substance, and may be calculated from the velocities of waves in the substance by means of the equations above, which give  $\psi$  in terms of  $\nu_m$ , and  $\nu_m$  in terms of  $v_1$  and  $v_2$ . The velocities  $v_1$  and  $v_2$  can be calculated from the bulk and rigidity moduli of elasticity. It is found that the specific heats of solids calculated in this way agree very well with those observed. In particular, at very low temperatures the specific heat is found to be proportional to the cube of the absolute temperature, and so becomes negligible at temperatures near zero.

The free energy of the solid can easily be calculated. We have seen that the free energy of a set of  $\mathcal{N}$  similar oscillators of frequency  $\nu$  is given by

$$F = k\mathcal{N}\theta \log(1 - e^{-h\nu/k\theta}) + \mathcal{N}\epsilon_0.$$

The free energy of the solid is the sum of the free energies corresponding to all the sets of vibrations in it, so that

$$F = E_0 + 4\pi h\theta \left( \frac{2}{v_1^3} + \frac{1}{v_2^3} \right) \int_0^{\nu_m} \nu^2 \log(1 - e^{-h\nu/k\theta}) d\nu,$$

where  $E_0$  is the energy in the solid when there is no vibration. As before, put  $x = h\nu/k\theta$  and  $h\nu_m = k\psi$ , so that

$$F = E_0 + \frac{9\mathcal{N}k\theta^4}{\psi^3} \int_0^{\psi/\theta} x^2 \log(1 - e^{-x}) dx.$$

\* For the method of evaluating the integral see section 11.

When  $\theta$  is very small, the integral becomes  $\int_0^\infty x^2 \log(1 - e^{-x}) dx = -\pi^4/45$ , so that

$$F = E_0 - \frac{\pi^4 \mathcal{N} k \theta^4}{5 \psi^3} \quad \dots \quad (9)$$

The entropy of the solid at low temperatures is then given by

$$\Phi = -\frac{\partial F}{\partial \theta} = \frac{4\pi^4 \mathcal{N} k \theta^3}{5 \psi^3} \quad \dots \quad (10)$$

According to this, when  $\theta = 0$ , then  $\Phi = 0$ , for any solid. But at  $\theta = 0$  all substances are solids, so that  $\Phi = 0$  at  $\theta = 0$  for all substances. This result is *Nernst's Heat Theorem*, sometimes called the *Third Law of Thermodynamics*.\*

Since  $\Phi = -\frac{\partial F}{\partial \theta}$ , we have  $\frac{\partial F}{\partial \theta} = 0$  at  $\theta = 0$ . The Gibbs-Helmholtz equation  $E = F - \theta \frac{\partial F}{\partial \theta}$  shows that  $E = F$  at  $\theta = 0$ , and also that  $\frac{\partial E}{\partial \theta} = 0$  at  $\theta = 0$ , so that at  $\theta = 0$ ,  $\frac{\partial F}{\partial \theta}$  and  $\frac{\partial E}{\partial \theta}$  are both zero.

## 11. Theory of Heat Radiation.

Instead of a solid cubical block we consider a hollow cube filled with radiation. The energy density of this radiation is independent of the nature of the walls and depends only on their temperature. If the walls are perfect reflectors, the radiation will form stationary trains of waves, and the number of possible frequencies between  $\nu$  and  $\nu + d\nu$  will be  $\frac{8\pi\nu^2 d\nu}{c^3}$ , where  $c$  is the velocity of light in a vacuum.

This follows, by differentiation with respect to  $\nu$ , from the expression  $\frac{8}{3}\pi(\nu/v)^3$  found in section 10 for the number of frequencies between 0 and  $\nu$  per unit volume. The additional factor 2 arises because the light waves are transverse, so that the radiation travelling in any direction may be regarded as made up of two parts polarized in two perpendicular directions, just as in the case of the transverse waves in the solid block previously considered. The energy in any one of the stationary vibrations in the box will be determined by the amplitude of the vibration just as in the case of the oscillators, so that according to the quantum theory the possible energies of the vibration will be given by  $\epsilon = n h \nu$  where  $n = 0, 1, 2, 3, \&c.$  The average energy of the vibrations having frequencies between  $\nu$  and  $\nu + d\nu$  will therefore be  $\frac{h\nu}{e^{h\nu/k\theta} - 1}$ , and the energy density  $E_\nu d\nu$  in these vibrations will be given by

$$E_\nu = \frac{8\pi\nu^3 h}{c^3 (e^{h\nu/k\theta} - 1)}.$$

\* Cf. J. K. Roberts, *Heat and Thermodynamics*, Chap. XVIII.

Here  $E_\nu$  may be defined as the energy density per unit range of frequency. This expression for  $E_\nu$  was first obtained by Planck by means of the quantum theory. It is found to agree well with experiments on the distribution of the energy in the spectrum of black body radiation. When  $h\nu/k\theta$  is large, the expression takes the form

$$E_\nu = \frac{8\pi\nu^3 h}{c^3} e^{-h\nu/k\theta},$$

known as Wien's formula. When  $h\nu/k\theta$  is very small,  $e^{h\nu/k\theta} = 1 + h\nu/k\theta$ , and Planck's expression becomes

$$E_\nu = \frac{8\pi\nu^2 k\theta}{c^3},$$

so that each vibration has energy  $k\theta$ , as in the case of the oscillators and solid body. According to Newtonian dynamics we should expect all values of the energy of a vibration or oscillator to be possible, since energy is supposed to be capable of continuous variation. The possible values of the energy of a vibration on the quantum theory are given by  $\epsilon = n h\nu$ ,  $n = 0, 1, 2, 3$ , &c., so that if  $h$  were indefinitely small all values of the energy would be possible since  $n$  can be as large as we please. The part of the quantum theory so far considered differs from the classical theory only in the supposition that  $h$  has a finite value instead of an indefinitely small value. If we put  $h = 0$  in Planck's formula for  $E_\nu$ , it becomes

$$E_\nu = \frac{8\pi\nu^2 k\theta}{c^3}.$$

This result is therefore that to which Newtonian dynamics leads, and it agrees with Planck's formula when  $h\nu/k\theta$  is very small, that is for low frequencies and high temperatures. The total energy density  $E$  in the radiation is given by

$$E = \int_0^\infty E_\nu d\nu.$$

If we put  $E_\nu = 8\pi\nu^2 k\theta/c^3$  we get an infinite value of  $E$ , which is of course impossible. The observed distribution of energy in the spectrum agrees with Planck's formula and differs entirely from that given by the classical theory even for frequencies of quite ordinary values. Newtonian dynamics therefore fails to explain heat radiation and so cannot be universally true as was formerly supposed. Of the other formulæ obtained in this chapter by means of the quantum theory, all those which do not contain  $h$  agree with Newtonian dynamics, while those containing  $h$  entirely disagree. The formulæ of the quantum theory all agree with experience, while those of the classical theory only do so when they agree with the quantum theory also.

If we put

$$E_\nu = \frac{8\pi\nu^3 h}{c^3(e^{h\nu/k\theta} - 1)} = \frac{8\pi k^3 \theta^3}{c^3 h^2} \frac{x^3}{e^x - 1},$$

we get

$$E = \frac{8\pi h}{c^3} \int_0^\infty \frac{\nu^3 d\nu}{e^{h\nu/k\theta} - 1} = \frac{8\pi k^4 \theta^4}{c^3 h^3} \int_0^\infty \frac{x^3 dx}{e^x - 1},$$

where  $x = h\nu/k\theta$ . Then

$$\begin{aligned} \int_0^\infty \frac{x^3 dx}{e^x - 1} &= \int_0^\infty (e^{-x} + e^{-2x} + e^{-3x} + \dots) x^3 dx \\ &= 6(1 + \frac{1}{2^4} + \frac{1}{3^4} + \dots) = \frac{\pi^4}{15}. \end{aligned}$$

Hence

$$E = \frac{8}{15} \frac{\pi^5 k^4 \theta^4}{c^3 h^3}.$$

Thus according to Planck's formula the total energy density of black

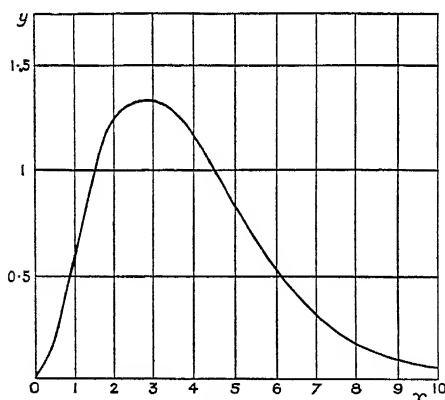


Fig. 2

body radiation in an enclosure is proportional to the fourth power of the absolute temperature, in agreement with Stefan's law. The relation between

$y = \frac{x^3}{e^x - 1}$  and  $x$  is shown in fig. 2. We see that as  $x$  increases  $y = \frac{x^3}{e^x - 1}$  in-

creases to a maximum value and then diminishes.  $E_\nu$  the energy density per unit range of frequency is equal to  $8\pi k^3 \theta^3 y / c^3 h^2$ , and so for

any given temperature  $E_\nu$  is proportional to  $y$ , and for any given value of  $y$   $E_\nu$  is proportional to the cube of the absolute temperature.

The energy density  $E_\lambda$  of the radiation per unit range of wavelength  $\lambda$  can be got by putting  $\nu = c/\lambda$  in  $E_\nu d\nu = \frac{8\pi h \nu^3 d\nu}{c^3(e^{h\nu/k\theta} - 1)}$ , and writing  $E_\nu d\nu = -E_\lambda d\lambda$ . This gives

$$E_\lambda = \frac{8\pi h c}{\lambda^5 (e^{hc/k\theta\lambda} - 1)}.$$

The wave-length  $\lambda_m$  for which  $E_\lambda$  is a maximum is got by putting  $dE_\lambda/d\lambda = 0$ . This gives

$$e^{hc/k\theta\lambda_m} = \frac{5}{5 - hc/k\theta\lambda_m},$$

the solution of which is  $\frac{hc}{k\theta\lambda_m} = 4.9651\dots$ . The two equations  $E = \frac{8\pi^5 k^{10}}{15c^3 h^3}$  and  $\frac{hc}{k\theta\lambda_m} = 4.9651$  enable the constants  $k$  and  $h$  to be calculated from the experimental values of  $E$  and  $\lambda_m$ . In this way it is found that  $h = 6.53 \times 10^{-27}$  erg-sec., and  $k = 1.37 \times 10^{-16}$  erg/degree. The most probable values of these constants are believed to be  $h = 6.554 \times 10^{-27}$  and  $k = 1.372 \times 10^{-16}$ . The energy  $h\nu$  is usually called a quantum of energy of frequency  $\nu$ . If  $\nu_m$  denotes the frequency of the light for which  $E_\lambda$  is a maximum so that  $c = \lambda_m \nu_m$  then we have  $h\nu_m = 4.9651 k\theta$ . The average kinetic energy of one molecule of a gas, as we have seen, is  $\frac{3}{2}k\theta$ , so that the quantum  $h\nu_m$  is equal to 3.31 times the average kinetic energy of a gas molecule at the same temperature. The fraction of the possible vibrations of frequency  $\nu$  which have energy  $n h\nu$  is equal to  $w_n = ae^{-nh\nu/k\theta}$  (cf. section 5; and  $\frac{1}{a} \sum e^{-nh\nu/k\theta} = 1/(1 - e^{-h\nu/k\theta})$ , or  $w_n = e^{-nh\nu/k\theta} (1 - e^{-h\nu/k\theta})$ ).

The fraction which have no energy is therefore  $w_0 = 1 - e^{-h\nu/k\theta}$ , so that the fraction which have any energy is  $e^{-h\nu/k\theta}$ . For example, at 0° C. the fraction of the vibrations of the frequency of yellow light ( $\nu = 5 \times 10^{14}$ ) which have any energy is only about  $e^{-87}$  or  $10^{-38}$ . The fraction of the vibrations of frequency  $\nu_m$  which have any energy is  $e^{-4.965}$  or about one in 140. According to classical dynamics the average energy of the vibrations of any frequency should be  $k\theta$ .

## 12. Einstein's Theory of Heat Radiation.

So far we have supposed that the energy is distributed among the parts of the system considered in such a way as to make the entropy or the number of possible microscopic states a maximum, but we have not considered the nature of the actions between the parts by which this equilibrium state is produced. An interesting way (due to Einstein) of getting Planck's formula throws light on this question, and will now be considered. Let a large number of molecules or atoms be contained in a perfectly reflecting enclosure so that there must be equilibrium between the black body radiation in the enclosure and the atoms. Also, let each atom be only capable of existing in one or other of a series of states, having energies  $\epsilon_1, \epsilon_2, \epsilon_3, \dots$ , and let the numbers of atoms in these states be  $N_1, N_2, N_3, \dots$ . Then the energy density of the radiation  $E_\nu$  per unit range of frequency  $\nu$  may be taken to be given by Wien's law,

$$E_\nu = \nu^3 f(0/\nu),$$

which can be deduced from purely thermodynamical considerations; and the

numbers of the atoms in the different states may be taken to be given by

$$N_1 = p_1 e^{-\epsilon_1/k\theta}, \quad N_2 = p_2 e^{-\epsilon_2/k\theta} \dots$$

The radiation of frequency  $\nu$  is supposed to be absorbed by atoms in the state having energy  $\epsilon_n$ , and as the result of this absorption these atoms are changed to another state having energy  $\epsilon_m$ . The number of atoms changed from state  $n$  to state  $m$  in this way will be proportional to  $N_n$  and to  $E_\nu$ , and so may be put equal to  $\alpha N_n E_\nu$ , where  $\alpha$  is a constant. The reverse change is supposed also to occur and in the same way the number changing from  $m$  to  $n$  may be put equal to  $\beta N_m E_\nu$ . The change from  $m$  to  $n$  is also supposed to occur spontaneously without any action of the radiation, and the number changing in this way may be put equal to  $\gamma N_m$ . In a state of equilibrium then we have

$$\alpha N_n E_\nu = \gamma N_m + \beta N_m E_\nu,$$

or

$$E_\nu (\alpha N_n - \beta N_m) = \gamma N_m.$$

Putting  $N_n = p_n e^{-\epsilon_n/k\theta}$ , and  $N_m = p_m e^{-\epsilon_m/k\theta}$ ,

we get

$$E_\nu (\alpha p_n e^{(\epsilon_m - \epsilon_n)/k\theta} - \beta p_m) = \gamma p_m.$$

At high temperatures  $E_\nu$  becomes very large, and  $\gamma p_m$  may be supposed independent of the temperature, so that we must have  $\alpha p_n - \beta p_m = 0$ ,

and therefore

$$E_\nu \alpha p_n (e^{(\epsilon_m - \epsilon_n)/k\theta} - 1) = \gamma p_m.$$

Now  $E_\nu$  only contains  $\theta$  in the function  $f(\theta/\nu)$  of  $\theta/\nu$ , so that  $(\epsilon_m - \epsilon_n)/k\theta$  must also be a function of  $\theta/\nu$ , which suggests that  $\epsilon_m - \epsilon_n$  should be proportional to  $\nu$ . Let  $\epsilon_m - \epsilon_n = h\nu$ , where  $h$  is a constant, so that

$$E_\nu = \frac{\gamma p_m / \alpha p_n}{e^{h\nu/k\theta} - 1}.$$

At high temperatures this gives

$$E_\nu = \frac{(\gamma p_m / \alpha p_n) k\theta}{h\nu}.$$

But at high temperatures the quantum theory and classical dynamics agree in giving

$$E_\nu = 8\pi\nu^2 k\theta / c^3,$$

so that we must have

$$\frac{(\gamma p_m / \alpha p_n)}{h\nu} = \frac{8\pi\nu^2}{c^3},$$

and therefore at any temperature

$$E_\nu = \frac{8\pi\nu^3 h}{c^3 (e^{h\nu/k\theta} - 1)},$$

which is Planck's formula.

The radiation of frequency  $\nu$  absorbed by the atoms in the above calculation must be equal to that emitted, so that when an atom goes from state  $m$  to state  $n$  we must suppose that it emits radiation of frequency  $\nu$  having energy  $\epsilon_m - \epsilon_n$ . According to this, when an atom emits an amount of energy, say  $\epsilon$ , in the form of radiation, the frequency of the radiation emitted is given by the equation  $\epsilon = h\nu$ . This result is the basis of Bohr's theory of spectral lines and was first put forward by him.

## BOHR'S THEORY OF SPECTRA

13. According to Bohr's theory an atom can exist only in a series of states having energies  $\epsilon_1, \epsilon_2, \epsilon_3 \dots$ , and when it changes from a state having energy  $\epsilon_m$  to another having less energy  $\epsilon_n$  the frequency of the radiation emitted is given by  $\epsilon_m - \epsilon_n = h\nu$ . The possible energies are determined in accordance with principles analogous to those of the quantum theory as outlined above. The application of Bohr's theory to the spectrum of atomic hydrogen, of which it gives a remarkably complete explanation, will now be considered. The frequencies of the lines in this spectrum are given with great accuracy by the equation

$$\nu = K \left( \frac{1}{n_1^2} - \frac{1}{n_2^2} \right),$$

where  $n_1 = 1, 2, \text{ or } 3$ ;  $n_2 = 2, 3, 4 \dots$ ; and  $K$  is a constant.

Bohr supposes that the hydrogen atom can only exist in one or other of a series of definite states having energies  $\epsilon_1, \epsilon_2, \epsilon_3 \dots$ , and that it only emits radiation when it jumps from one of those states to another with less energy. The frequency of the radiation emitted is then given by the equation  $h\nu = \epsilon_{n_2} - \epsilon_{n_1}$ , where  $\epsilon_{n_2}$  is the energy of the atom in the first state and  $\epsilon_{n_1}$  that in the second state. If we suppose that  $\epsilon_{n_2} = A - Kh/n_2^2$ , and  $\epsilon_{n_1} = A - Kh/n_1^2$ , where  $A$  is another constant, then we get the above equation for  $\nu$ .

The hydrogen atom is believed to consist of a minute nucleus having a charge of positive electricity  $e$ , and an electron with the charge  $-e$ . The mass  $m$  of the electron is only  $1/1860$  of that of the nucleus, and the electron describes an orbit round the nucleus. Let the position of the electron be determined by rectangular co-ordinates,  $x$  and  $y$ , with the nucleus at the origin. The kinetic energy of the electron  $T = \frac{1}{2}m(\dot{x}^2 + \dot{y}^2)$ , and the potential energy  $V = A - e^2/r$ , where  $r = \sqrt{x^2 + y^2}$ . The volume in the state space enclosed by a surface of constant total energy  $\epsilon = T + V$  is

$$m^2 \int dx dy d\dot{x} d\dot{y},$$

where  $\int d\dot{x} d\dot{y}$  for a given value of  $T$  is equal to the area of a circle of radius  $\sqrt{2T/m}$ , so that the above integral is equal to

$$2\pi m \int T dx dy,$$

or

$$2\pi m \int (\epsilon - A + e^2/r) 2\pi r dr.$$

The greatest value of  $r$  occurs when  $T = 0$ , and so is equal to  $e^2/(A - \epsilon)$ ,

so that we get  $4\pi^2 m \int_0^{e^2/(A-\epsilon)} \{(\epsilon - A)r + e^2\} dr = \frac{2\pi^2 m e^4}{A - \epsilon}$

for the volume enclosed by the surface of constant energy  $\epsilon$ .

If we now suppose that

$$m^2 \int dx dy d\dot{x} d\dot{y} = n^2 h^2 \quad (n = 0, 1, 2, 3, \dots),$$

we get

$$\frac{2\pi^2 m e^4}{A - \epsilon_n} = n^2 h^2,$$

or

$$\epsilon_n = A - \frac{2\pi^2 m e^4}{n^2 h^2}.$$

The frequencies of the light emitted are then given by

$$h\nu = \epsilon_{n_2} - \epsilon_{n_1} = \frac{2\pi^2 m e^4}{h^2} \left( \frac{1}{n_1^2} - \frac{1}{n_2^2} \right).$$

This result agrees with the equation

$$\nu = K \left( \frac{1}{n_1^2} - \frac{1}{n_2^2} \right)$$

if  $K = \frac{2\pi^2 m e^4}{h^3}$ .

On substituting the known values of  $e/m$ ,  $c$ , and  $h$  we obtain  $K = 3.294 \times 10^{15}$ , which agrees as well as could be expected with the value  $3.290 \times 10^{15}$ , which is deduced from the frequencies of the lines in the hydrogen spectrum.

The spectrum of ionized helium can be explained in the same way. The helium atom is believed to consist of a nucleus with a positive charge  $2e$ , and two electrons describing orbits round the nucleus. Ionized helium consists of helium atoms which have lost one electron. The potential energy of the ionized helium atom is therefore

$$V = A - 2e^2/r,$$

so that we get

$$\epsilon_n = A - \frac{8\pi^2 m e^4}{n^2 h^2},$$

and

$$\nu = \frac{8\pi^2 m e^4}{h^3} \left( \frac{1}{n_1^2} - \frac{1}{n_2^2} \right) = 4K \left( \frac{1}{n_1^2} - \frac{1}{n_2^2} \right).$$

The frequencies in the spectrum of ionized helium agree with this expression. The corresponding formula for the frequencies emitted by an atom with a nucleus having a positive charge  $Ne$  and one electron is

$$\nu = \frac{2\pi^2 N^2 m e^4}{h^3} \left( \frac{1}{n_1^2} - \frac{1}{n_2^2} \right) = N^2 K \left( \frac{1}{n_1^2} - \frac{1}{n_2^2} \right).$$

Some spectral lines due to doubly ionized lithium, for which  $N = 3$ , have been observed.

In the above calculations of  $\nu$  we supposed the nucleus to be at rest at the origin. This is not quite correct, since the nucleus although much heavier than the electron has not an infinite mass. The small

correction required can be easily found. The centre of mass of the atom may be supposed at rest at the origin, so that the kinetic energy is  $\frac{1}{2}mv^2 + \frac{1}{2}M\left(\frac{mv}{M}\right)^2$ , where  $m$  is the mass of the electron, and  $M$  that of the nucleus. Thus  $T = \frac{1}{2}mv^2\left(1 + \frac{m}{M}\right)$ . The potential energy is  $V = A - \frac{Ne^2M}{r(m+M)}$ , since  $r\left(1 + \frac{m}{M}\right)$  is the distance from the nucleus to the electron. To correct for the mass of the nucleus it is therefore only necessary to replace  $m$  by  $m\left(1 + \frac{m}{M}\right)$ , and  $e^2$  by  $e^2\left(1 + \frac{m}{M}\right)^{-1}$ . The corrected expression for the frequencies is therefore

$$\nu = \frac{2\pi^2 N^2 m e^4}{h^3} \frac{M}{M+m} \left( \frac{1}{n_1^2} - \frac{1}{n_2^2} \right),$$

or

$$\nu = \frac{N^2 K M}{M+m} \left( \frac{1}{n_1^2} - \frac{1}{n_2^2} \right).$$

The mass of the helium nucleus is 4/1.008 or 3.97 times that of the hydrogen nucleus, so that if  $\nu_H$  is the frequency of a hydrogen line and  $\nu_{He}$  that of an ionized helium line having the same values of  $n_1$  and  $n_2$ , then

$$\frac{\nu_{He}}{\nu_H} = 4 + 2.97 \frac{m}{M}.$$

Accurate measurements of  $\nu_{He}/\nu_H$  by Fowler give for this ratio 4.001638, so that

$$M/m = 2.97/0.001638 = 1813.$$

This result agrees very well with the ratio of the mass of an hydrogen atom to that of an electron, as determined by other methods, and so gives an interesting confirmation of the theory.

It is interesting to compare the results of Bohr's theory with those of classical dynamics. In the case of the hydrogen atom we should expect the fundamental frequency of the radiation emitted to be equal to that of the revolution of the electron round the nucleus, and harmonics having multiples of this frequency might also be expected. The frequency of revolution would not be constant, but would increase as the atom gradually lost energy.

The theory of central orbits shows that the electron describes an ellipse with the nucleus at one focus, and the energy of the atom is equal to  $A - e^2/2a$ , where  $A$  is a constant and  $a$  is the length of the major axis of the orbit. The frequency of revolution ( $\omega$ ) is equal to  $\frac{e}{2\pi m^{1/2} a^{3/2}}$ . The energy of the atom on the quantum theory is

$A - Kh/n^2$ , so that we may write  $a_n = \frac{n^2 e^2}{2Kh}$ , where  $a_n$  is the major axis of the orbit in the  $n$ th possible state. The frequency  $\omega_n$  in the  $n$ th state is therefore given by

$$\omega = \frac{2K}{n^3},$$

since  $K = 2\pi^2 me^4/h^3$ . Bohr's theory gives

$$\nu = K \left( \frac{1}{n_1^2} - \frac{1}{n_2^2} \right),$$

which may be written  $\nu = 2K \int_{n_1}^{n_2} \frac{dx}{x^3}$ ,

where  $x$  may be defined by  $E = A - Kh/x^2$ , so that the possible states of Bohr's theory are got by putting  $x = 1, 2, 3, \dots$ . Thus on Bohr's theory  $x$  can only have one or other of these values, whereas on the classical theory it can vary continuously. The frequency of revolution  $\omega$  is equal to  $2K/x^3$ , so that the average value of  $\omega$  as  $x$  changes from  $n_2$  to  $n_1$  may be taken to be given by

$$\bar{\omega} = \frac{2K}{n_2 - n_1} \int_{n_1}^{n_2} \frac{dx}{x^3} = \frac{\nu}{n_2 - n_1}.$$

Hence  $\nu = (n_2 - n_1)\bar{\omega}$ , or the frequencies emitted on Bohr's theory are multiples of the average frequency of revolution of the electron. When  $n_1$  and  $n_2$  are very large and nearly equal numbers,  $\omega_{n_2}$  will be nearly equal to  $\omega_{n_1}$  and  $\bar{\omega}$  will be nearly equal to  $\omega$ , and we shall have

$$\nu = (n_2 - n_1)\omega,$$

so that in this case the Bohr theory and the classical theory agree approximately. In this case  $\nu$  is small, so that in the theory of spectra as in that of heat radiation the classical and quantum theories agree for very low frequencies.

#### 14. Cases with more than one Quantum Number.

In the cases so far considered the possible energies ( $\epsilon_1, \epsilon_2, \epsilon_3 \dots$ ) of the systems have been determined by finding the volume in the state space corresponding to a constant value of the energy, and then assuming that the volumes ( $V_1, V_2 \dots$ ) enclosed by the surfaces corresponding to the possible values of the energy were multiples of some power of Planck's constant  $h$ . In the case of monatomic gas atoms we assumed  $V_n = nh^3$ , in the case of simple harmonic vibrations  $V_n = nh$ , and in the case of atoms having a nucleus and one electron  $V_n = n^2 h^2$ . In these cases it was possible to express the possible energies in terms of a single integer ( $n = 0, 1, 2, 3 \dots$ ) or quantum number.

The quantum theory has been applied successfully to several more complicated cases in which the possible energies are determined by two or more independent quantum numbers. In such cases the volume of the surfaces of constant energy in the state space cannot be used to determine the possible energies.

Consider a periodic atomic system determined by co-ordinates  $q_1, q_2 \dots$  and the corresponding momenta  $p_1, p_2 \dots$ . The representative point in the state space describes a curve which in the case of a periodic system is closed. The projection of this closed curve on one of the co-ordinate planes ( $p'q'$ ) will also be a closed curve, the area of which may be represented by  $\int p'dq'$ . In certain cases it has been found possible to obtain correct results by supposing that

$$\int p_1 dq_1 = n_1 h, \quad \int p_2 dq_2 = n_2 h, \quad \int p_3 dq_3 = n_3 h,$$

where  $n_1, n_2, n_3 \dots$  are independent integers  $0, 1, 2, 3 \dots$ . As a simple example, consider the case of a particle of mass  $m$  vibrating in the  $x, y$  plane under the action of a force directed towards the origin and equal to  $\mu r$  where  $r = \sqrt{x^2 + y^2}$ . The particle describes elliptical orbits with period  $2\pi\sqrt{m/\mu}$ . The energy  $E$  is given by

$$E = \frac{1}{2}m(\dot{x}^2 + \dot{y}^2) + \frac{1}{2}\mu r^2.$$

Let

$$E_1 = \frac{1}{2}m\dot{x}^2 + \frac{1}{2}\mu x^2,$$

so that

$$\frac{(m\dot{x})^2}{2mE_1} + \frac{\mu x^2}{2E_1} = 1.$$

Thus for a given value of  $E_1$  the representative point describes an ellipse of area  $2\pi E_1 \sqrt{m/\mu}$  on the  $(x, m\dot{x})$  plane. Assume then

$$2\pi E_1 \sqrt{m/\mu} = n_1 h \quad (n_1 = 0, 1, 2 \dots).$$

In the same way let  $\frac{1}{2}m\dot{y}^2 + \frac{1}{2}\mu y^2 = E_2$ , so that

$$2\pi E_2 \sqrt{m/\mu} = n_2 h \quad (n_2 = 0, 1, 2, 3 \dots).$$

Then

$$E = E_1 + E_2 = \frac{h}{2\pi\sqrt{m/\mu}} (n_1 + n_2).$$

But the frequency of vibration  $\omega$  is equal to  $(2\pi\sqrt{m/\mu})^{-1}$ , so that

$$E = (n_1 + n_2) \hbar \omega = n \hbar \omega \quad (n = 0, 1, 2, 3 \dots).$$

This result agrees with that obtained for the one-dimensional oscillator, as it should, since the vibrations along  $x$  are independent of those along  $y$ . The two-dimensional oscillator can be regarded as equivalent to the two independent one-dimensional oscillators. If we calculate

the volume in the state space corresponding to constant energy  $E$  it is easily found to be  $E^2/2\omega^2$ . If we put this equal to  $n\hbar^2$ , we get

$$E = (2n)^{1/2}\hbar\omega.$$

This gives for the frequency  $\nu$  of the radiation emitted on Bohr's theory

$$\nu = \omega\sqrt{2}(\sqrt{n_1} - \sqrt{n_2}).$$

This result seems improbable, since we should expect the radiation emitted to have a frequency equal to  $\omega$ . If we put the volume equal to  $n^2\hbar^2/2$  we get  $E = n\hbar\omega$ , and so

$$\nu = (n_1 - n_2)\omega,$$

which agrees with  $\nu = \omega$ , provided  $n_1 - n_2 = 1$ .

### 15. Sommerfeld's Theory of the Fine Structure of Spectral Lines.

An important contribution to the quantum theory of spectra has been made by Sommerfeld. So far, we have regarded the mass  $m$  of the electron as independent of its velocity, but this is not exactly true, since it is known that the mass varies with the velocity according to the law

$$m = \frac{m_0}{\sqrt{1 - v^2/c^2}},$$

where  $m_0$  is the mass when the velocity  $v$  is zero, and  $c$  is the velocity of light. Sommerfeld examined the effect on Bohr's theory of allowing for this variation of the mass. Consider an atom consisting of a nucleus with positive charge  $Ne$  and an electron with charge  $-e$ . The kinetic energy of the electron is given by

$$T = m_0c^2 \left\{ \frac{1}{\sqrt{1 - \beta^2}} - 1 \right\},$$

where  $\beta = v/c$ . The constant angular momentum of the electron,  $p$ , is equal to  $mr^2\dot{\theta}$ , so that

$$p = \frac{m_0r^2\dot{\theta}}{\sqrt{1 - \beta^2}}, \quad \dots \dots \dots (11)$$

where  $r$  is the distance of the electron from the nucleus, and  $\theta$  the angle between  $r$  and a fixed line through the nucleus in the plane of the orbit. The potential energy,  $V$ , of the electron is equal to  $A - Ne^2/r$ , so that the total energy  $E$  is given by

$$E = m_0c^2 \left\{ \frac{1}{\sqrt{1 - \beta^2}} - 1 \right\} + A - \frac{Ne^2}{r}. \quad \dots \dots \dots (12)$$

As before, we suppose that the atom emits no radiation except when the electron jumps from one of its possible orbits to another one with less energy. We have  $v^2 = \dot{r}^2 + r^2\dot{\theta}^2$ , which with (11) on putting  $r = 1/u$  gives

$$\left( \frac{du}{d\theta} \right)^2 + u^2 = \frac{m_0^2c^2\beta^2}{p^2(1 - \beta^2)}.$$

Equation (12) with this gives

$$\left(\frac{du}{d\theta}\right)^2 + u^2 = \frac{m_0^2 c^2}{p^2} \left\{ \left(1 + \frac{E - A + Ne^2 u}{m_0 c^2}\right)^2 - 1 \right\}.$$

The solution of this equation is

$$u = C(1 + \alpha \cos \gamma \theta),$$

where

$$\gamma^2 = 1 - \left(\frac{Ne^2}{pc}\right)^2, \quad \dots \dots \dots (13)$$

$$C = \frac{Ne^2}{\gamma^2 p^2} \left(m_0 + \frac{E - A}{c^2}\right),$$

$$\alpha^2 = 1 + \frac{\gamma^2 (E - A)(E - A + 2m_0 c^2)}{(1 - \gamma^2)(E - A + m_0 c^2)^2}. \quad \dots \dots \dots (14)$$

Thus if  $\gamma$  had the value 1 the orbit would be an ellipse, and if  $\gamma$  is nearly equal to 1 we may say that the orbit is an ellipse with its major axis slowly rotating about the origin in the plane of the orbit. When  $\gamma\theta = \pi, 3\pi, 5\pi \dots$ , so that  $\cos \gamma\theta = -1$ , then  $r = u^{-1}$  has its maximum value and  $\theta = \pi/\gamma, 3\pi/\gamma, 5\pi/\gamma \dots$ , so that the major axis turns through an angle  $2\pi \left(\frac{1-\gamma}{\gamma}\right)$  during each revolution of the electron. The two co-ordinates  $r$  and  $\theta$  and the corresponding momenta  $m\dot{r}$  and  $m r^2 \dot{\theta}$  will be the co-ordinates of the representative point in the state space. We assume that

$$\int m r^2 \dot{\theta} d\theta = n_1 \hbar \quad (n_1 = 0, 1, 2, 3 \dots),$$

and

$$\int m \dot{r} dr = n_2 \hbar \quad (n_2 = 0, 1, 2, 3 \dots).$$

In this way we obtain two quantum numbers to define the possible states of the atom. The first integral is taken over one complete revolution so that, since  $m r^2 \dot{\theta} = p$  is constant, we get

$$\int_0^{2\pi} p d\theta = 2\pi p = n_1 \hbar.$$

This fixes the possible values of the angular momentum  $p$ . The second integral is taken from one minimum value of  $r$  to the next one, or from  $\theta = 0$  to  $\theta = 2\pi/\gamma$ .

We have  $\int m \dot{r} dr = -p \int \frac{du}{d\theta} dr = -p \int_0^{2\pi/\gamma} \frac{du}{d\theta} \frac{dr}{d\theta} d\theta$ .

Substituting for  $\frac{du}{d\theta}$  and  $\frac{dr}{d\theta}$  from  $u = C(1 + \alpha \cos \gamma \theta)$ , and integrating, we find

$$\frac{1}{\sqrt{1 - \alpha^2}} - 1 = \frac{n_2 \hbar}{2\pi \sqrt{p^2 - (Ne^2/c)^2}}. \quad \dots \dots \dots (15)$$

This equation determines the possible values of the eccentricity  $\alpha$  of the orbit for each possible value of  $p$ . The possible values of the energy of the atom  $E$  can now be obtained from equations (13), (14), and (15). The term  $Ne^2/c$  in (15) is small compared with  $p$  so that for a first approximation it may be neglected. Thus, putting  $p = n_1 \hbar / 2\pi$ , equation (15) becomes

$$1 - \alpha^2 = \frac{n_1^2}{(n_1 + n_2)^2}.$$

In (14) for a first approximation we may put  $\gamma^2 = 1$  and neglect  $E - A$  in comparison with  $m_0 c^2$ , so that with (13) it becomes

$$\alpha^2 - 1 = \frac{(E - A) h^2 n_1^2}{2\pi^2 m_0 N^2 e^4}.$$

Hence

$$E = A - \frac{2\pi^2 m_0 N^2 e^4}{h^2 (n_1 + n_2)^2}.$$

The frequency of the light emitted is then given by  $h\nu = E_{n'} - E_n$ , so that

$$\nu = \frac{2\pi^2 m_0 N^2 e^4}{h^3} \left( \frac{1}{n^2} - \frac{1}{n'^2} \right)$$

where  $n$  has been put for  $n_1 + n_2$ . This is the same result as was obtained before when the mass of the electron was assumed constant, so that the approximations just made are equivalent to neglecting the variation of  $m$ .

The equation  $1 - \alpha^2 = \frac{n_1^2}{(n_1 + n_2)^2}$  gives the possible eccentricities of the orbits, or if  $b$  and  $a$  denote the minor and major semi-axes of an orbit, since  $\frac{b}{a} = \sqrt{1 - \alpha^2}$ , we have

$$\frac{b}{a} = \frac{n_1}{n_1 + n_2}.$$

Thus, for example, if  $n_1 + n_2 = 4$  then  $b/a$  may be equal to  $1/4$ ,  $2/4$ ,  $3/4$ , or  $4/4$ , but all these orbits correspond to nearly the same value of the energy  $E$ , since this is determined by  $n_1 + n_2 = n$  when  $m$  is assumed constant. These orbits also have equal major axes for, when  $E - A$  is small compared with  $m_0 c^2$ , and  $\gamma = 1$ , we have  $\frac{1}{C} = \frac{b^2}{a}$ ,

so that

$$a = \frac{a^2}{b^2 C},$$

or

$$a = \left( \frac{n}{n_1} \right)^2 \frac{1}{C} = \frac{n^2 h^2}{4\pi^2 N m_0 e^2},$$

which shows that  $a$  is determined by  $n = n_1 + n_2$ . To obtain the effect of the variation of  $m$  it is necessary to calculate  $E$  more accurately.

Equation (14), if we retain only the first power of  $(E - A)/m_0 c^2$ , gives

$$E - A = m_0 c^2 (k + \frac{3}{2} k^2), \quad \dots \dots \dots (16)$$

where

$$k = \frac{(\alpha^2 - 1)(1 - \gamma^2)}{2\gamma^2}.$$

Equations (13) and (15) give

$$k = - \frac{n_0^2}{2(n_1 + n_2)^2} \left( 1 + \frac{n_0^2}{n_1(n_1 + n_2)} \right),$$

where  $n_0 = \frac{2\pi N e^2}{h c}$ , and squares and higher powers of  $\frac{n_0}{n_1}$  have been left out. Substituting this value of  $k$  in (16) and putting  $n = n_1 + n_2$ , we get

$$E = A - \frac{m_0 c^2 n_0^2}{2n^2} \left\{ 1 + \frac{n_0^2}{n^2} \left( \frac{1}{4} + \frac{n_2}{n_1} \right) \right\},$$

or, since

$$K = \frac{2\pi^2 m_0 e^4}{h^3} \quad \text{and} \quad n_0 = \frac{2\pi N e^2}{hc},$$

$$E = A - \frac{N^2 K h}{n^2} \left\{ 1 + \frac{2N^2 K h}{m_0 c^2 n^2} \left( \frac{1}{4} + \frac{n_2}{n_1} \right) \right\}.$$

Thus it appears that the effect of the variation of  $m$  with the velocity is to subtract a very small quantity  $\frac{2N^4 K^2 h^2}{m_0 c^2 n^4} \left( \frac{1}{4} + \frac{n_2}{n_1} \right)$  from the expression for the energy of the atom. If the atom jumps from a state having quantum numbers  $n_1'$  and  $n_2'$  to another having quantum numbers  $n_1$  and  $n_2$ , then the frequency emitted is given by

$$\nu = N^2 K \left( \frac{1}{n_2^2} - \frac{1}{n_1'^2} \right) + \frac{2N^4 K^2 h}{m_0 c^2} \left\{ \frac{1}{n_1^4} \left( \frac{1}{4} + \frac{n_2}{n_1} \right) - \frac{1}{n_1'^4} \left( \frac{1}{4} + \frac{n_2'}{n_1'} \right) \right\},$$

where  $n = n_1 + n_2$  and  $n' = n_1' + n_2'$ .

The Balmer series of lines in the spectrum of atomic hydrogen is got by putting  $N = 1$ ,  $n = 2$ , and  $n' = 3, 4, 5, \dots$ . The first line  $H_\alpha$  corresponds to  $n = 2$  and  $n' = 3$ . For this line  $n_1$  therefore may be 1 or 2, and  $n_2$  may be 1 or 0, so that  $\frac{1}{n_1^4} \left( \frac{1}{4} + \frac{n_2}{n_1} \right)$  may be

$$1/16 (1/4 + 0/2) = 1/64,$$

or

$$1/16 (1/4 + 1/1) = 5/64.$$

In the same way  $n_2'$  may be 0, 1, or 2, and  $n_1'$  may be 3, 2, or 1. The values  $n_1 = 0$  and  $n_1' = 0$  are not considered to be possible values, since they correspond to a line orbit meeting the nucleus. They also make  $\frac{n_2'}{n_1'}$  and  $\frac{n_2}{n_1}$  infinite, and so are clearly not allowable. The three possible values of  $\frac{1}{n_1'^4} \left( \frac{1}{4} + \frac{n_2'}{n_1'} \right)$  for  $n' = 3$  are therefore

$$1/81 (1/4 + 0/3) = 1/324,$$

$$1/81 (1/4 + 1/2) = 3/324,$$

$$1/81 (1/4 + 2/1) = 9/324.$$

We should expect each of these three values to give a spectral line with each of the two values of  $\frac{1}{n_1^4} \left( \frac{1}{4} + \frac{n_2}{n_1} \right)$ , so that the  $H_\alpha$  line should have six components.

The three values of  $\frac{1}{n_1'^4} \left( \frac{1}{4} + \frac{n_2'}{n_1'} \right)$  are nearly equal, so that we should expect the six components to form two groups of three lines each, the difference between each line in one group and the corresponding one in the other being

$$\frac{2K^2 h}{m_0 c^2} \left( \frac{5}{64} - \frac{1}{64} \right) = \frac{K^2 h}{8m_0 c^2}.$$

Actually, the  $H_\alpha$  line is found to be a doublet for which the difference between the two frequencies is nearly equal to  $\frac{K^2 h}{8m_0 c^2} = 1.09 \times 10^{10}$  vibrations per second.

This is in agreement with the theory, for the three components of each group are so close together that they cannot be resolved into separate lines. The fine structure of the other Balmer series lines, and also that of several lines in the spectrum of ionized helium, for which  $N = 2$ , is found in the same way to agree

very well with that predicted by this theory. For example, the helium line 4686 is found to have six components the positions of which agree remarkably well with those calculated.

### 16. The Zeeman Effect. Explanation on Classical Theory.

In 1896 Zeeman discovered that spectral lines are affected when the source of light is put in a strong magnetic field. When the light is examined along the direction of the field each line becomes a doublet, but if it is viewed at right angles to the field a triplet of equally spaced lines is seen. The middle line of the triplet coincides with the single line emitted when there is no magnetic field, and the other two are in the same positions as the doublet lines seen along the field. The light of the doublet lines is circularly polarized in opposite directions, and the triplet lines are plane polarized, the middle one perpendicular to the field and the other two parallel to it.

The change of frequency  $\nu$  is given by

$$\Delta\nu = \frac{eH}{4\pi m} = 1.41 \times 10^6 H \text{ vibrations per second.}$$

Thus if  $H = 10,000$ , then  $\Delta\nu = 1.41 \times 10^{10}$ , so that, if  $\nu = 3 \times 10^{14}$ , the frequency and so the wave-length is changed by about one in 20,000. The normal Zeeman effect just described occurs only with simple single lines; doublets and triplets, however close, give more complex effects. It is found, however, that lines which give a complex Zeeman effect in ordinary magnetic fields may nevertheless give a normal effect in very strong fields. This is called the Paschen-Back effect.

The normal effect can be explained on the classical theory if we suppose that the light is emitted by an electron vibrating about a position of equilibrium towards which it is attracted by a force proportional to its displacement.

The motion of the electron may then be resolved into three mutually perpendicular simple harmonic motions all having the same frequency. If one of these is taken to be along the magnetic field its frequency will not be altered by the field, and the light due to it at right angles to the field will be plane polarized perpendicular to the field, while along the field it will give no light. The two perpendicular simple harmonic motions in the plane perpendicular to the field may be represented by

$$\begin{aligned} x &= a \cos pt, \\ y &= b \sin pt, \end{aligned}$$

so that the orbit is an ellipse, since

$$\frac{x^2}{a^2} + \frac{y^2}{b^2} = \cos^2 pt + \sin^2 pt = 1.$$

Let  $x = x_1 + x_2$  and  $y = y_1 + y_2$ , where

$$\begin{aligned}x_1 &= c_1 \cos pt, & x_2 &= c_2 \cos pt, \\y_1 &= c_1 \sin pt, & y_2 &= -c_2 \sin pt,\end{aligned}$$

so that  $a = c_1 + c_2$ , and  $b = c_1 - c_2$ . Then  $x_1^2 + y_1^2 = c_1^2$ , and  $x_2^2 + y_2^2 = c_2^2$ . Thus we see that the motion round the ellipse can be regarded as made up of two circular motions in opposite directions. These circular motions in the plane perpendicular to the magnetic field will be affected by the field. If  $H$  denotes the strength of the magnetic field, and  $\mu$  the restoring force at unit distance from the equilibrium position, then we have

$$mr\omega^2 = \mu r + Her\omega,$$

where  $r$  is the radius of the circle,  $m$  is the mass of the electron,  $\omega$  its angular velocity, and  $e$  its charge.

Hence, if  $\omega_0$  is the value of  $\omega$  when  $H = 0$ ,

$$m(\omega^2 - \omega_0^2) = Hew,$$

or

$$2m\omega\Delta\omega = Hew.$$

But  $\omega = 2\pi\nu$ , so that  $\Delta\nu = \frac{He}{4\pi m}$ .

If we change  $\omega$  to  $-\omega$  we get  $\Delta\nu = -\frac{He}{4\pi m}$ . Thus we see that along the field we shall get two lines circularly polarized in opposite directions having frequencies equal to  $\nu_0 \pm \frac{He}{4\pi m}$ .

The classical theory therefore gives a satisfactory explanation of the normal Zeeman effect. Since, however, we have good reason for believing in the nucleus theory of atoms this theory cannot be accepted.

### 17. The Zeeman Effect. Explanation on Quantum Theory.

The Zeeman effect can also be explained on the quantum theory, as was shown by Sommerfeld. Consider a hydrogen atom and take axes  $x, r, \theta$  with the nucleus at the origin. Let the magnetic field  $H$  be along the  $x$  axis, and let  $r$  be the distance of the electron from the  $x$  axis, and  $\theta$  the angle between  $r$  and a fixed plane containing the  $x$  axis. The equations of motion of the electron are then

$$\begin{aligned}m\ddot{x} &= \frac{-e^2x}{(r^2 + x^2)^{3/2}}, \\ \frac{d}{dt}(mr^2\dot{\theta}) &= Her\dot{r}, \\ m\ddot{r} &= \frac{-e^2r}{(r^2 + x^2)^{3/2}} - Her\dot{\theta} + mr\dot{\theta}^2.\end{aligned}$$

The motion along  $x$  is unaffected by the magnetic field. We can show that if we change to new axes rotating about  $x$  with angular velocity  $\omega = He/2m$ , then the motion of the electron referred to the rotating axes will be practically the same as the motion relative to the fixed axes when  $H = 0$ . To change to the new axes we put  $\theta + \omega t$  for  $\theta$ , and also introduce into the equations of motion the apparent forces due to the rotation. These forces are the centrifugal force  $mr\omega^2$  and the Coriolis force, the components of which are  $2mr\dot{\omega}$  perpendicular to  $r$  in the plane perpendicular to  $x$ , and  $2mr\omega\dot{\theta}$  along  $r$ . The equations of motion referred to the new axes are therefore

$$\frac{d}{dt}(mr^2\dot{\theta}) = Her\dot{r} - 2mr\dot{\omega},$$

$$m\ddot{r} = \frac{-e^2r}{(r^2 + x^2)^{3/2}} - Her\dot{\theta} - Her\omega + 2mr\omega\dot{\theta} + mr\dot{\theta}^2 + mr\omega^2.$$

If  $\omega = He/2m$  the first of these equations becomes  $\frac{d}{dt}(mr^2\dot{\theta}) = 0$ , which is the corresponding equation relative to fixed axes when  $H = 0$ . The second equation becomes

$$m\ddot{r} = \frac{-e^2r}{(r^2 + x^2)^{3/2}} - \frac{rH^2e^2}{4m} + mr\dot{\theta}^2.$$

The term  $\frac{rH^2e^2}{4m}$  can be neglected in ordinary cases, so that this equation also is practically the same as for fixed axes with  $H = 0$ . We may therefore suppose that the possible orbits of the electron on the quantum theory are the same referred to the rotating axes as those referred to fixed axes with  $H = 0$ . The possible values of  $p = mr^2\dot{\theta}$  are therefore  $n\hbar/2\pi$  ( $n = 0, 1, 2 \dots$ ). The change in the energy due to the rotation with the axes is

$$\delta E = \delta(\frac{1}{2}mr^2\dot{\theta}^2) = mr^2\dot{\theta}\delta\dot{\theta} = p\delta\dot{\theta}.$$

But  $\delta\dot{\theta} = \omega$ , so that

$$\delta E = p\omega = \frac{n\hbar}{2\pi} \frac{He}{2m}.$$

The change in the frequency  $\nu$  of the light emitted is therefore given by

$$\delta\nu = \frac{He}{4\pi m} (n - n').$$

Here  $n - n' = 0, \pm 1, \pm 2, \pm 3 \dots$ . According to this the light emitted would give a line of the unaltered frequency corresponding to  $n - n' = 0$  and a series of equally spaced lines on either side of this one, with the frequency differences  $He/4\pi m$ . The observed light in the normal Zeeman effect shows the central unaltered line ( $n - n' = 0$ ) and

one on each side of it ( $n - n' = \pm 1$ ). In some cases very faint lines corresponding to  $n - n' = \pm 2$  and  $n - n' = \pm 3$  have been observed.

### 18. The Stark Effect.

In 1913 Stark discovered that the spectral lines emitted by an atom are modified when the atom is in a strong electric field. This effect is especially well marked in the case of atomic hydrogen and helium. The lines are split up into a number of components the distances between which are proportional to the strength of the field. The electric field alters the electron orbits and so changes the possible energies of the atom. In the case of atomic hydrogen and ionized helium each line given by the equation

$$\nu = N^2 K \left( \frac{1}{n^2} - \frac{1}{n'^2} \right)$$

consists of a group of lines very near together. The small differences between the frequencies in these groups are due to the slightly different effects of the variation of the mass of the electron with its velocity on the energies of the differently shaped possible orbits corresponding to any one value of  $n$ . When the atom is placed in an electric field the energy associated with any orbit is changed, and the change is different for differently shaped orbits. Consequently, the field increases the separation between the lines in the group of lines and may also cause some of them to split into two or more lines. The quantum theory of this effect has been worked out by Epstein and others, the results of the theory agreeing in a most remarkable way with what is observed. The theory, however, is too long and complicated for discussion in this book.

Applications of the quantum theory to photo-electricity, thermionics, X-rays, and other branches of physics are discussed in the chapters dealing with these subjects.

### *Quantum Theory and Wave Theory.*

19. The quantum theory is based on two assumptions: (1) atoms can exist only in one or other of a series of possible states having definite energies  $\epsilon_1, \epsilon_2, \epsilon_3 \dots$ ; (2) when an atom emits radiation the frequency  $\nu$  of the radiation is given by

$$h\nu = \epsilon - \epsilon',$$

where  $\epsilon$  is the energy of the atom before the emission and  $\epsilon'$  that after the emission. It is also believed that when radiation is absorbed by an atom the amount absorbed is equal to  $h\nu$ . When X-rays are produced by the impact of electrons on a solid body then the greatest frequency  $\nu_m$  of the X-rays is given by

$$h\nu_m = \epsilon,$$

where  $\epsilon$  is the kinetic energy of the electron. Also, when electrons are set free from matter by X-rays or light of frequency  $\nu$ , the energy given to each electron by the radiation is equal to  $h\nu$ , whatever may be the intensity of the radiation. In general, whenever radiation of frequency  $\nu$  is absorbed or emitted by matter, the energy absorbed

or emitted seems to be a multiple of  $h\nu$ . An exception to this rule occurs in the scattering of X-rays by the electrons in light elements, in which case, according to Compton, the electron gets energy  $\epsilon$  less than  $h\nu$  and the frequency  $\nu'$  of the scattered radiation is given by

$$\epsilon = h\nu - h\nu'.$$

This, however, is not really an exception, for we may suppose energy  $h\nu$  absorbed and then  $h\nu'$  emitted. These results seem to show that radiation consists of elements or quanta each having energy  $h\nu$ . According to the wave theory, radiation consists of transverse waves, in which the energy is distributed continuously and not in finite quanta. It is difficult to see how the two theories can be reconciled. Consider, for example, the interference method by which Michelson has recently estimated the diameters of certain stars. The light from the star is received on two mirrors 15 ft. apart in the plane of the waves, and the two reflected beams are brought together in an eye-piece and produce interference bands. According to the wave theory this shows that the individual wave fronts extend unbroken over areas at least 15 ft. across, so that at any instant the light disturbances at two points in the plane of the waves are equal even when the two points are 15 ft. apart. Also any desired fraction of the light may be cut off by putting up a screen with an aperture of any desired size in it. If the light consists of quanta which cannot be divided without altering the frequency, then we must suppose that they are small, so that the number passing through an aperture is proportional to the area of the aperture. But Michelson's experiment just mentioned shows that the wave fronts are at least 15 ft. across. It is clear, therefore, that many quanta must exist in the area of one continuous wave front. Also interference can be obtained with homogeneous light with a path difference of at least 100 cm., so that the train of waves in the light must be at least 100 cm. long without any break. Such facts on the wave theory show that there can be trains of waves occupying a volume of several hundred cubic feet, and such trains can be divided into many parts without altering the wave-length. Interference phenomena are not affected by changing the intensity of the light. An interesting experiment was tried by G. I. Taylor to test this. He photographed an interference pattern with a short exposure in strong light and also with an exposure of several hundred hours in very weak light. The two photographs were identical in appearance. The phenomena of interference, diffraction, and polarization support the wave theory so strongly that it is difficult to believe that it is based on an erroneous conception of reality, but on the other hand the phenomena of emission and absorption involving the quantum  $h\nu$  seem to support the quantum theory so strongly that it also must have some basis of truth behind it.

When an interference pattern is produced on a screen by two beams of light the energy is concentrated into the bright bands, as may be proved by means of a linear thermopile. If the light consists of small quanta each having energy  $h\nu$ , we must then suppose that these quanta only fall on the screen on the bright bands. The quanta of one beam must therefore be deviated from their straight paths by those of the other beam. The paths of the quanta must be along the direction of the flow of energy. If a quantum having energy  $\epsilon$  is travelling in the  $x$  direction with the velocity of light  $c$ , and performing oscillations of some kind of frequency  $\nu$ , then, since the momentum of the quantum is  $\epsilon/c$  in the  $x$  direction, we may suppose that

$$\int_0^\lambda \frac{\epsilon}{c} dx = h, \quad \text{or} \quad \frac{\epsilon\lambda}{c} = h,$$

so that  $\epsilon = h\nu$ , since  $\nu = c/\lambda$ . Here  $\lambda$  is the distance the quantum moves along  $x$  during one complete vibration. According to this, the quantum itself conforms to the quantum theory rule,

$$\int p dq = n h, \quad \text{with } n = 1.$$

When light is reflected from a moving mirror its wave-length is changed, as it should be according to the wave theory. This effect can also be explained on the quantum theory. Suppose a quantum having energy  $h\nu$  and momentum  $h\nu/c$  to be reflected normally from a mirror moving with velocity  $v$  in the opposite direction to the quantum. Let the reflected quantum have energy  $h\nu'$  and momentum  $h\nu'/c$ . During the collision between the quantum and the mirror let the force exerted on the quantum by the mirror be  $F$ , and let the duration of the collision be  $\tau$ . Then we have by the conservation of momentum

$$\frac{h\nu}{c} + \frac{h\nu'}{c} = \int_0^\tau F dt,$$

and by the conservation of energy

$$h\nu' - h\nu = v \int_0^\tau F dt,$$

so that

$$\frac{\nu' - \nu}{\nu' + \nu} = \frac{v}{c},$$

or

$$\frac{\nu'}{\nu} = \frac{c + v}{c - v},$$

which is the same result as is given by the wave theory.

A quantum theory of the diffraction grating has been suggested by Duane and Compton. Suppose a quantum of energy  $h\nu$  to be incident on a plane transmission grating at an angle  $\phi$  and diffracted at an

angle  $\theta$ . The momentum parallel to the surface of the grating lost by the quantum is

$$\frac{h\nu}{c} (\sin \phi + \sin \theta).$$

This must be equal to the momentum in the same direction received by the grating, which we assume equal to  $\frac{n h \nu'}{c}$ , where  $n = 0, \pm 1, \pm 2 \dots$ , and  $\nu' = c/d$ ,  $d$  being the grating space, so that  $c/d$  may be regarded as the frequency of the grating. Hence

$$\sin \phi + \sin \theta = \frac{nc}{\nu d},$$

which agrees with the usual grating formula. It must be admitted that this theory of the grating is not very convincing.

## 20. Bohr's Correspondence Principle.

An important principle due to Bohr is the "Correspondence Principle". We have seen in several cases that the results of the quantum theory agree with those of classical dynamics when the frequency considered is very small. In particular, the frequency of the light emitted by an atom containing an electron is equal to the frequency of revolution of the electron, when the energy emitted is very small. A similar result can be shown to be generally true. Consider a system determined by co-ordinates  $q_1, q_2 \dots$  and momenta  $p_1, p_2 \dots$ , the latter being defined as usual by  $p_s = \frac{\partial E}{\partial \dot{q}_s}$ , where  $E$  denotes the energy of the system expressed as a function of the  $q$ 's and  $\dot{q}$ 's. If the kinetic energy  $T$  is a homogeneous quadratic function of the  $\dot{q}$ 's, we have by Euler's theorem

$$2T = \Sigma \dot{q} \frac{\partial T}{\partial \dot{q}} = \Sigma \dot{q} \frac{\partial E}{\partial \dot{q}} = \Sigma p \dot{q}.$$

Let the system be periodic so that all the  $q$ 's and  $p$ 's repeat their values at equal intervals  $\tau$ , and let  $I = \int_0^\tau \Sigma p \dot{q} dt$ , so that  $I/\tau$  is twice the average value of the kinetic energy during one period. Now suppose that the energy of the system is changed slightly from  $E$  to  $E + \delta E$ , so that  $I$  changes by  $\delta I$ , where

$$\delta I = \int_0^\tau \Sigma (\dot{q} \delta p + p \delta \dot{q}) dt + (\Sigma p \dot{q})_{t=\tau} \delta \tau.$$

The last term represents the variation due to the change in  $\tau$ .

Now

$$\int_0^\tau p \delta \dot{q} dt = (p \delta q)_{t=\tau} - \int_0^\tau \delta q \dot{p} dt,$$

since  $\delta \dot{q} dt = d\delta q$ , and we suppose the time reckoned from the same value

of  $q$  in the varied and unvaried motions, so that  $\delta q = 0$  when  $t = 0$ .

$$\text{Hence} \quad \delta I = \int_0^\tau \Sigma (q \delta p - p \delta q) dt + \{ \Sigma p (\delta q + \dot{q} \delta \tau) \}_{t=\tau}.$$

But  $q$  at  $t = 0$  is equal to  $q$  at  $t = \tau$  since the motion is periodic; hence, when  $t = \tau$ ,

$$\delta q + \dot{q} \delta \tau = 0.$$

$$\text{We have therefore} \quad \delta I = \int_0^\tau \Sigma (\dot{q} \delta p - \dot{p} \delta q) dt.$$

Now if  $E$  is expressed as a function of the  $p$ 's and  $q$ 's we have

$$\dot{p} = -\frac{\partial E}{\partial q} \text{ and } \dot{q} = \frac{\partial E}{\partial p},$$

$$\text{so that} \quad \delta I = \int_0^\tau \Sigma \left( \frac{\partial E}{\partial p} \delta p + \frac{\partial E}{\partial q} \delta q \right) dt = \int_0^\tau \delta E dt.$$

But  $\delta E$  is constant so that  $\delta I = \tau \delta E$ , or, if  $\omega$  denotes the frequency or  $\tau^{-1}$ ,

$$\omega = \frac{\delta E}{\delta I} = \frac{dE}{dI}.$$

According to Bohr's theory we have  $h\nu = \epsilon - \epsilon'$ , which we may write

$$\nu = \frac{\Delta \epsilon}{h}.$$

Also if  $\int p_1 dq_1 = n_1 h$ ,  $\int p_2 dq_2 = n_2 h$ , and so on, we may put

$$I = \int_0^\tau \Sigma p \dot{q} dt = \Sigma n h = n h \quad (n = 0, 1, 2, \dots),$$

so that when  $\epsilon$  changes from  $\epsilon$  to  $\epsilon'$  and  $I$  changes from  $I$  to  $I'$  we shall have

$$I - I' = \Delta I = (n - n')h$$

and

$$\nu = (n - n') \frac{\Delta \epsilon}{\Delta I}.$$

If  $n$  and  $n'$  are very large integers and  $n - n'$  is small, we may suppose

$$\Delta \epsilon = \delta \epsilon \text{ and } \Delta I = \delta I, \text{ so that, since } \omega = \frac{\delta \epsilon}{\delta I}, \text{ we get } \nu = (n - n') \omega.$$

Thus according to the quantum theory when  $\epsilon - \epsilon'$  is very small the frequency emitted is a multiple of the frequency of oscillation of the system, in agreement with classical theory. In any case we may put

$$\bar{\omega} = \frac{1}{\Delta I} \int_r^I \frac{d\epsilon}{dI} dI = \frac{\Delta \epsilon}{\Delta I},$$

so that  $\bar{\omega}$  is an average value of the frequency of the system during the change of the energy from  $\epsilon$  to  $\epsilon'$ . Hence we have

$$\nu = (n - n')\bar{\omega},$$

and we may say that the frequencies emitted on the quantum theory are multiples of the average frequency of the emitting system. In the case of Balmer's series,

$$\nu = K \left( \frac{1}{n'^2} - \frac{1}{n^2} \right),$$

where  $n' = 2$  and  $n = 3, 4, 5 \dots$ , frequencies of lines are given by

$$\nu = \bar{\omega}_{32} \times 1, \quad \nu = \bar{\omega}_{42} \times 2, \quad \nu = \bar{\omega}_{52} \times 3,$$

and so on. Here, for example,  $\bar{\omega}_{42}$  is the average frequency of revolution of the electron during the transition from the state given by  $n = 4$  to that given by  $n = 2$ . According to this, lines in a series like the Balmer series may be said to correspond to the harmonics of the frequency of the atom which would be emitted on the classical theory, if the atom had a fixed frequency of vibration. According to Bohr, results given by the classical theory as to the different lines emitted may be supposed to apply to the corresponding lines predicted by the quantum theory. This is Bohr's Correspondence Principle. For example, a simple harmonic oscillator of frequency  $\omega$  on the classical theory only emits radiation of frequency  $\omega$ . But the quantum theory gives for the energies in the possible states

$$\epsilon_n = n\hbar\omega \quad (n = 0, 1, 2, 3 \dots),$$

so that the frequencies emitted should be given by

$$h\nu = \epsilon_n - \epsilon_{n'},$$

or

$$\nu = (n - n')\omega.$$

The Correspondence Principle therefore indicates that  $n - n' = 1$ , or that the intensities of the harmonics of the fundamental frequency are zero. In the case of the hydrogen atom we might assume the electron to be describing a particular orbit with constant frequency  $\omega$ , and then calculate the relative intensities of the lines having frequencies  $\omega, 2\omega, 3\omega \dots$  to be expected on the classical theory. The results obtained might be expected to give the relative intensities of the corresponding lines of the quantum theory. Thus the line  $\nu = K(\frac{1}{4} - \frac{1}{16}) = \bar{\omega}_{42} \times 2$  would be expected to have the intensity of the first harmonic of the radiation from an electron describing an orbit with constant frequency  $\bar{\omega}_{42}$ . The Correspondence Principle has proved of great value. It enables the intensity and state of polarization of lines to be predicted correctly in many cases.

## 21. Adiabatic Invariants.

As we have seen, the possible states of a periodic atomic system can be determined in certain cases by means of the equations

$$\int p_1 dq_1 = n_1 h, \quad \int p_2 dq_2 = n_2 h, \quad \&c.$$

The quantity  $I = \int \Sigma p \dot{q} dt$  is therefore also equal to  $n h$  ( $n = 0, 1, 2 \dots$ ). Also, since the kinetic energy  $T$  is given by

$$2T = \Sigma p \dot{q},$$

we have for a periodic system

$$2\bar{T} = \frac{1}{\tau} \int_0^\tau \Sigma p \dot{q} dt = \frac{n h}{\tau},$$

where  $\bar{T}$  denotes the average value of  $T$  during one complete oscillation of period  $\tau$ ; or, since  $\tau = \nu^{-1}$ , we get

$$\frac{2\bar{T}}{\nu} = n h.$$

Suppose now that the atomic system is acted on by some external field of force such as an electric or magnetic field, and that this field is initially zero and gradually increases in intensity to a final constant value. The field will alter the motion of the system and will change the energies of the possible states. In order that the quantum condition  $\int p dq = n h$  may apply in the presence of the field it is clear that  $\int p dq$  must remain unchanged as the field is applied. Quantities which remain unchanged in value according to classical dynamics when a field is gradually applied are called "adiabatic invariants", following Einstein, Ehrenfest, and Burgers, to whom this idea is due. It can be shown on the classical theory that  $2\bar{T}/\nu$  is an adiabatic invariant, and it is suggested that only adiabatic invariants should be put equal to multiples of  $h$  for the purpose of determining the possible states of a periodic system on the quantum theory. The discovery of an adiabatic invariant of an atomic system may therefore enable its possible states to be determined by the quantum theory.

## 22. Physical Interpretation of the Quantum Theory.

The physical basis of the quantum theory has not yet been discovered. We do not know why atomic systems should only be able to exist in certain definite states in which they do not radiate, and we do not know why the frequency of the radiation emitted should be given by  $h\nu = \varepsilon - \varepsilon'$ . The most promising attempt to devise a model of an atom to illustrate the quantum theory is due to Whittaker, and we will now consider a modification of his model.

Suppose an atom consists of a filament in the form of a closed curve which is a perfect conductor of magnetism. We may suppose the filament to consist of a mixture of equal quantities of north and south magnetism which can move freely along the filament. Suppose now that an electron with a charge  $e$  is moving towards this filament. The number of lines of electric induction coming out of the electron is  $4\pi e$ , so that the number passing through the filament is  $e\omega$ ,  $\omega$  being the solid angle subtended by the filament at the electron. The electric displacement current through the filament due to the motion of the electron is therefore  $e\dot{\omega}/4\pi$ . The work to take a unit pole once round the filament or the magnetomotive force due to this current is  $e\dot{\omega}$ , so that a magnetic current will be induced in the filament by the motion of the electron. This magnetic current produces an electric field which retards the motion of the electron.

The electric potential due to a constant magnetic current  $i$  is equal to  $i\omega$ , so that the force on the electron is  $-ie\frac{d\omega}{ds}$ , where  $ds$  is an element of the path of the electron. Hence, if  $E$  denotes the kinetic energy of the electron, we have

$$E = E_0 - e \int i \frac{d\omega}{ds} ds,$$

where  $E_0$  is the initial value of  $E$ . Let the number of lines of electric induction due to the magnetic current  $i$  be  $KSi$ , where  $S$  is a constant and  $K$  the specific inductive capacity of the surrounding space. Then

$$KS \frac{di}{dt} - e \frac{d\omega}{dt} = 0,$$

since we suppose the filament to be a perfect conductor of magnetism. Hence

$$KS i = e\omega$$

if we take  $i = 0$  when  $\omega = 0$ . We therefore have

$$E = E_0 - \frac{e^2}{KS} \int \omega \frac{d\omega}{ds} ds,$$

or

$$E = E_0 - \frac{e^2 \omega^2}{2KS}.$$

As the electron approaches the filament  $\omega$  increases and so  $E$  diminishes, so that if  $E_0$  is not too big the electron may be brought to rest when

$$E_0 = \frac{e^2 \omega^2}{2KS}.$$

If the electron does not pass through the filament it will be repelled and will move away until  $\omega = 0$  and  $E = E_0$ , so that it will get back its initial kinetic energy. If the electron has enough kinetic energy it may go through the filament and move off to an infinite distance after going through. In this case  $\omega$  changes by  $4\pi$  so that the final energy of the electron is given by  $E = E_0 - \frac{8\pi^2 e^2}{KS}$ . The electron therefore loses energy  $8\pi^2 e^2 / KS$  when it goes right through, whatever its initial energy may be, but if it does not go right through it loses no energy. The filament, therefore, can have energy zero or energy  $8\pi^2 e^2 / KS$ . If a second electron goes through the filament in the opposite direction to the first one it will get the energy  $8\pi^2 e^2 / KS$  from the filament. If, however, it approaches the

filament in the same direction as the first one the energy of the filament will be increased again. In this case we have

$$KS\dot{i} - e\omega = \text{constant},$$

and, when  $\omega = 0$ ,  $KS\dot{i} = 4\pi e$ , so that  $KS\dot{i} - e\omega = 4\pi e$ . Hence

$$E = E_0 - \frac{e^2}{KS} \left( 4\pi\omega + \frac{\omega^2}{2} \right).$$

Thus, if the second electron goes right through, its final energy will be given by

$$E = E_0 - \frac{1}{2} \frac{(4\pi e)^2}{KS},$$

and in the same way for a third electron

$$E = E_0 - \frac{5}{2} \frac{(4\pi e)^2}{KS},$$

and so on. The energy lost by the electrons is gained by the filament as the energy of the electric field of the filament. The possible values of the energy of the filament are therefore

- (1) 0,
- (2)  $\frac{1}{2} \frac{(4\pi e)^2}{KS},$
- (3)  $\frac{4}{2} \frac{(4\pi e)^2}{KS},$
- (4)  $\frac{9}{2} \frac{(4\pi e)^2}{KS},$

and so on. The possible energies of the filament are therefore given by

$$E_n = \frac{n^2}{2} \frac{(4\pi e)^2}{KS} \quad (n = 0, 1, 2, 3 \dots).$$

Thus the filament can exist in a series of states having definite energies, and it can change from one state to the next one by giving up the energy difference to an electron. In order that an atom may emit radiation, some kind of oscillations must take place in it, and we shall suppose that this oscillation is analogous to the oscillatory discharge of a condenser through a circuit having self induction. In such oscillations the energy of the system changes from electric to magnetic energy and back again. If  $C$  is the capacity of the condenser and  $Q$  the charge on one plate, then

$$E = \frac{1}{2} \frac{Q^2}{C} + \frac{1}{2} L \left( \frac{dQ}{dt} \right)^2,$$

where  $E$  is the energy of the system and  $L$  the self induction of the circuit. The current is equal to  $-dQ/dt$ . The periodic time is given by

$$T = 2\pi\sqrt{LC}.$$

In the case of the filament we may suppose that it is a conductor for electricity as well as for magnetism, so that if the magnetic current in it is  $i$  and the electric current  $j$  then the energy  $E$  is given by

$$E = \frac{KS\dot{i}^2}{2} + \frac{1}{2} S\mu j^2,$$

since  $S\mu$  will be the self induction of the filament,  $\mu$  being the permeability, and  $K$  the specific inductive capacity of the surrounding space. Now the number of lines of electric induction passing through the filament is  $KSi = 4\pi ne$ , which would come out of a charge  $Q = \frac{KS i}{4\pi} = ne$ . We may suppose the electric field of the magnetic current to be due to two charges  $Q$  and  $-Q$  suitably distributed, and that the electric current  $j$  is equal to  $-dQ/dt$ . Hence we get

$$E = \frac{(4\pi Q)^2}{2KS} + \frac{1}{2}S\mu \left(\frac{dQ}{dt}\right)^2.$$

Comparing this with the corresponding equation for a condenser discharging through a circuit we conclude that the periodic time of the filament will be

$$T = 2\pi \sqrt{\frac{S\mu KS}{(4\pi)^2}} = \frac{S}{2} \sqrt{\mu K}.$$

The frequency is therefore  $\eta = \frac{2}{S\sqrt{\mu K}}$ , and so is independent of the energy  $E$ .

The radiation emitted according to the classical theory will have frequencies  $\eta, 2\eta, 3\eta \dots$ .

According to the quantum theory the frequencies emitted will be given by

$$E_n - E_{n'} = h\nu,$$

or

$$\frac{(4\pi e)^2}{2KS} (n^2 - n'^2) = h\nu,$$

so that the possible values of  $\nu$  will be multiples of  $\frac{(4\pi e)^2}{2KS\hbar}$ . The two theories therefore will agree if

$$\eta = \frac{2}{S\sqrt{\mu K}} = \frac{(4\pi e)^2}{2KS\hbar},$$

or if

$$\hbar = 4(\pi e)^2 \sqrt{\frac{\mu}{K}}.$$

As a matter of fact  $\hbar$  is nearly equal to  $2\pi(4\pi e)^2 \sqrt{\frac{\mu}{K}}$ , which is  $8\pi$  times greater than the value just obtained. The model, therefore, does not give the correct value of  $\hbar$ . It seems probable that atoms are really more complicated structures than is usually supposed, and that the physical interpretation of the quantum theory will not be clear until we know much more about them than we do now.

## THEORIES OF HEISENBERG, DE BROGLIE, AND SCHRÖDINGER

23. An important step in the direction of a real physical interpretation of the quantum theory has been recently initiated by Heisenberg and developed by Born and Dirac. De Broglie and Schrödinger have also attacked the problem in another new way which, however, turns out to be formally equivalent to that of Heisenberg. Schrödinger's development of De Broglie's ideas will be briefly outlined here, since it seems clear that he has really got at the root of the problem.

## 24. Fermat's Law and Hamilton's Principle. Least Action.

It is necessary to begin by discussing some preliminary questions. In geometrical optics the path of a ray of light may be determined by Fermat's law, according to which the time taken by the light to pass from one point  $A$  to another  $B$  along its actual path is equal to the time along any other path between the same two points which is very close to the actual path. If  $ds$  denotes an element of the path and  $u$  the velocity of the light along  $ds$ , then according to Fermat's law

$$\delta \int_A^B \frac{ds}{u} = 0,$$

since  $ds/u$  is the time taken to traverse the distance  $ds$ . The symbol  $\delta$  here indicates a very small change from the actual path to the other path, and the equation indicates that the change in the time from  $A$  to  $B$  when the path is changed from the actual path to any other path, very near to the actual path, is zero.

Fermat's law can be easily deduced from the wave theory of light. Consider a wave from  $A$  moving towards  $B$ . According to Huygens' principle we may suppose that secondary wavelets start from every point on the wave front at any instant. The effect at  $B$  due to a small area on the wave front will be very small unless the secondary waves from the different parts of this small area all arrive at  $B$  at the same instant. This means that the time to go from any point on the small area to  $B$  must be the same if the actual path of the ray from  $A$  to  $B$  passes through the small area. Thus as the wave moves from  $A$  to  $B$  a small area on it moves along the ray path, and at any point on the path the time for the light to go to  $B$  from all points on the small area is the same. It is clear therefore that the times from  $A$  to  $B$  along the actual path and all the paths very near to it are equal.

In the dynamical theory of any material system we have a principle which may be expressed in a similar way to Fermat's law in geometrical optics. This principle, known after its discoverer as Hamilton's principle, was in fact suggested to him by Fermat's law.

Consider a material system consisting of any number of particles having masses  $m_1, m_2, m_3, \dots$ . Let the co-ordinates of the particles be  $x_1, y_1, z_1; x_2, y_2, z_2; x_3, y_3, z_3, \dots$ ; and let  $t$  denote the time. Also suppose that the particles are moving in a field of force excited by the particles so that the system has potential energy as well as kinetic energy. We shall suppose that the total energy of the system is constant and that the potential energy depends only on the positions of the particles and not on their velocities.

The kinetic energy  $T$  of the system is then given by

$$2T = \Sigma m(\dot{x}^2 + \dot{y}^2 + \dot{z}^2),$$

and the potential energy  $V$  is a function of all the co-ordinates of the particles. It is convenient to abbreviate  $\Sigma m(\dot{x}^2 + \dot{y}^2 + \dot{z}^2)$  into  $\Sigma m\dot{x}^2$ , it being understood that the sign  $\Sigma$  then indicates a summation of all products like  $m\dot{x}^2$ , including the  $m\dot{y}^2$  and  $m\dot{z}^2$ .

We may regard the positions of the particles in the system as determined by the position of a single point the co-ordinates of which are  $x_1, y_1, z_1, x_2, y_2, z_2, x_3, y_3, z_3, \dots$ , that is, we suppose this representative point to be moving in an imaginary space of  $3N$  dimensions,  $N$  being the number of particles in our system. As the particles move, the representative point moves along a path in this imaginary space. Now consider two points  $A$  and  $B$  in this space on a path of the representative point. Let  $t_A$  be the time at which the point is at  $A$  and  $t_B$  the time at which it reaches  $B$ . Then according to Hamilton's principle

$$\delta \int_{t_A}^{t_B} (T - V) dt = 0,$$

where the  $\delta$  indicates a small change from the actual path from  $A$  to  $B$  to any other path from  $A$  to  $B$  which is very close to the actual path. The small change is not supposed to involve a change in the time  $t$ . That is, we suppose that to each point on the actual path there is a corresponding point on the varied path and that the time is the same at these two points.

Since

$$T = \frac{1}{2} \Sigma m \dot{x}^2,$$

we have

$$\delta T = \Sigma m \dot{x} \delta \dot{x}.$$

Also, since the  $x$  component of the force on a particle is equal to  $m\ddot{x}$ , we have

$$-\delta V = \Sigma (m \ddot{x} \delta x + m \ddot{y} \delta y + m \ddot{z} \delta z),$$

or, in the abbreviated form,

$$-\delta V = \Sigma m \ddot{x} \delta x.$$

Hence

$$\delta T - \delta V = \Sigma m \dot{x} \delta \dot{x} + \Sigma m \ddot{x} \delta x.$$

Now

$$\frac{d}{dt} (m \dot{x} \delta x) = m \ddot{x} \delta x + m \dot{x} \delta \dot{x},$$

so that

$$\delta T - \delta V = \frac{d}{dt} \Sigma m \dot{x} \delta x.$$

Therefore, since we are supposing that  $\delta(dt) = 0$ ,

$$\delta \int_{t_A}^{t_B} (T - V) dt = \int_{t_A}^{t_B} (\delta T - \delta V) dt = \int_{t_A}^{t_B} \frac{d}{dt} (\Sigma m \dot{x} \delta x) dt.$$

Hence

$$\delta \int_{t_A}^{t_B} (T - V) dt = \left[ \Sigma m \dot{x} \delta x \right]_{t_A}^{t_B}.$$

But the two paths are supposed to coincide at  $A$  and  $B$ , so that every  $\delta x$ ,  $\delta y$ , and  $\delta z$  is zero at  $A$  and  $B$ , and therefore

$$\delta \int_{t_A}^{t_B} (T - V) dt = 0.$$

Let the total energy be  $E = T + V$  so that  $T - V = 2T - E$ , and we have

$$\delta \int_{t_A}^{t_B} (T - V) dt = \delta \int_{t_A}^{t_B} 2T dt - \delta \int_{t_A}^{t_B} E dt.$$

But  $E$  is constant, so that

$$\int_{t_A}^{t_B} E dt = E(t_B - t_A),$$

and therefore, since  $\delta t = 0$ ,

$$\delta \int_{t_A}^{t_B} E dt = 0,$$

so that

$$\delta \int_{t_A}^{t_B} 2T dt = 0.$$

This result follows immediately from Hamilton's principle and is called the principle of least action.  $\int_{t_A}^{t_B} 2T dt$  is called the action.

If we suppose all the particles in the system fixed except one, then we have only to consider the three co-ordinates of this particle, and the path of the representative point is simply the path of the particle. Now consider the path of a ray of light between two points  $A$  and  $B$ , and the path of a particle between the same two points. For the path of the ray we have

$$\delta \int_A^B \frac{ds}{u} = 0,$$

and for the path of the particle, since  $2T = mv^2$  and  $v dt = ds$ ,

$$\delta \int_A^B m v ds = 0.$$

We see that the two paths will be identical if  $mv = k/u$ , where  $k$  is any constant.

## 25. Schrödinger's Wave Mechanics.

De Broglie suggested that a train of waves is associated with a moving electron, and Schrödinger goes one step farther and regards an electron as actually consisting of a train of waves.

The photo-electric effect and the scattering of X-rays by electrons suggest that light has the essential properties of material particles. It behaves as though it consisted of particles having energy  $h\nu$  and momentum  $h\nu/c$ . Again, the deflection of light by the gravitational field of the sun is the same as that of a particle moving with the velocity of light. An electromagnetic field has energy and momentum and

moves through space by growing in front and fading away behind, a process similar to wave motion.

The fact that light behaves like particles becomes comprehensible if particles are trains of waves like light. In the equation  $mv = k/u$  the velocity of the particle is  $v$  and the velocity of the train of waves which would go along the same path is  $u$ . If the particle is a train of waves, then the wave velocity must be  $u$  and the velocity of the train or group of waves must be  $v$ . Now it is well known that the velocity of a group of waves travelling in a dispersive medium is different from the wave velocity. We see, therefore, that if  $u$  is the wave velocity and  $v$  the group velocity of the waves which are the electron, then its motion will agree with Fermat's law and also with Hamilton's principle.

The relation between the wave velocity and the group velocity can be easily obtained. Consider two trains of waves of equal amplitude travelling in the same direction, and let the resultant amplitude be

$$\begin{aligned} y &= \cos 2\pi\nu \left( t - \frac{x}{u} \right) + \cos 2\pi\nu' \left( t - \frac{x}{u'} \right) \\ &= 2 \cos \pi \left( (\nu + \nu')t - \left( \frac{\nu}{u} + \frac{\nu'}{u'} \right) x \right) \cos \pi \left( (\nu - \nu')t - \left( \frac{\nu}{u} - \frac{\nu'}{u'} \right) x \right), \end{aligned}$$

or, putting  $\nu - \nu' = \delta\nu$  and  $\frac{\nu}{u} - \frac{\nu'}{u'} = \delta \left( \frac{\nu}{u} \right)$ ,

$$y = 2 \cos 2\pi\nu \left( t - \frac{x}{u} \right) \cos \pi \left( t\delta\nu - x\delta \left( \frac{\nu}{u} \right) \right).$$

Thus we get a train of waves of frequency  $\nu$  and wave velocity  $u$  with amplitude varying very slowly along  $x$  between 2 and  $-2$ .

The waves between two points at which the amplitude is zero form a group of waves, and the group travels with the velocity of its middle point, where the amplitude is 2. This group velocity  $v$  is given by

$$t\delta\nu - x\delta \left( \frac{\nu}{u} \right) = 0, \text{ so that } \frac{1}{v} = \frac{t}{x} = \frac{\delta \left( \frac{\nu}{u} \right)}{\delta\nu}.$$

The velocity  $v$  of the particle and the wave velocity  $u$  must therefore satisfy this equation.

Fermat's law gives the path of a ray of light and the laws of geometrical optics may be easily deduced from it. The phenomena of interference and diffraction which appear when the fine structure of the light distribution is examined do not follow Fermat's law. This law is merely an approximation valid so long as the scale of the phenomena considered is large compared with the wave-length of the light.

Hamilton's principle gives correct results for the motion of dynamical systems consisting of large bodies, but classical dynamics fails

to give a satisfactory account of atomic phenomena which are on a very small scale. Schrödinger points out the analogy between the failure of Fermat's law for small-scale optical phenomena and the failure of Hamilton's principle for atomic phenomena. According to his view both failures are due to the same cause, namely, that the scale of the phenomena considered is not large compared with the wave-lengths involved, so that the theories based on the paths of rays are inadequate and wave theory is required to give a satisfactory account of the phenomena.

## 26. Schrödinger's Wave Mechanics and the Hydrogen Atom.

Let us now consider a system consisting of one particle moving in a fixed field of force. If  $x$ ,  $y$ , and  $z$  are the co-ordinates of the particle, the kinetic energy  $T = \frac{1}{2}m(\dot{x}^2 + \dot{y}^2 + \dot{z}^2)$ , and we may take the potential energy  $V$  to be a function of  $x$ ,  $y$ , and  $z$  only.

Let  $W$  be a quantity defined by the equation

$$W = S(x, y, z) - Et,$$

where  $E = T + V$  is the total energy,  $t$  the time, and  $S$  a function of  $x$ ,  $y$ ,  $z$  such that  $\frac{\partial S}{\partial x} = m\dot{x}$ ,  $\frac{\partial S}{\partial y} = m\dot{y}$ , and  $\frac{\partial S}{\partial z} = m\dot{z}$ . We have then, since  $W$  is a function of  $t$ ,  $x$ ,  $y$ , and  $z$  only,

$$\begin{aligned} \frac{dW}{dt} &= \frac{\partial W}{\partial t} + \frac{\partial W}{\partial x} \dot{x} + \frac{\partial W}{\partial y} \dot{y} + \frac{\partial W}{\partial z} \dot{z} \\ &= -E + m\dot{x}^2 + m\dot{y}^2 + m\dot{z}^2 = 2T - (T + V) \\ &= T - V. \end{aligned}$$

Hence  $W = \int (T - V) dt + \text{constant}.$

Now suppose a family of surfaces drawn in the field of force so that  $W$  is constant over each surface, and increases from one surface to the next one by a constant amount  $\Delta W$ . The equation of a surface is then

$$W_0 + m\Delta W = S(x, y, z) - Et,$$

where  $m$  is equal to 0, 1, 2, 3, 4, &c. The gradient of  $W$  is a vector with components  $\partial W/\partial x$ ,  $\partial W/\partial y$ ,  $\partial W/\partial z$ , and so is perpendicular to the surfaces on which  $W$  is constant. If then  $s$  is the distance measured along a vector line of grad  $W$ , i.e. a line whose direction at any point is that of the vector, grad  $W$ , we have

$$\left(\frac{\partial W}{\partial s}\right)^2 = \left(\frac{\partial W}{\partial x}\right)^2 + \left(\frac{\partial W}{\partial y}\right)^2 + \left(\frac{\partial W}{\partial z}\right)^2,$$

so that  $\left(\frac{\partial W}{\partial s}\right)^2 = m^2(\dot{x}^2 + \dot{y}^2 + \dot{z}^2)$ , or  $\frac{\partial W}{\partial s} = mv.$

As the time increases, the surfaces on which  $W$  is constant move along with a velocity  $u$  given by

$$dW = 0 = \frac{\partial W}{\partial s} ds + \frac{\partial W}{\partial t} dt,$$

or, since  $ds = udt$ ,

$$\frac{\partial W}{\partial s} u + \frac{\partial W}{\partial t} = 0,$$

so that

$$u = E / \frac{\partial W}{\partial s} = \frac{E}{mv}.$$

Any path of the particle is given by  $\delta W = 0$ , according to Hamilton's principle, and is therefore a vector line of grad  $W$ , that is a line everywhere perpendicular to the surfaces on which  $W$  is constant. For if we consider such a line between two points  $A$  and  $B$ , then it is clear that  $W_B - W_A = \int_A^B (T - V) dt$  has the same value when the integration is taken along that line or along any other line between  $A$  and  $B$  very close to it, because  $W$  is unchanged for any small displacement normal to a vector line of grad  $W$ . The surfaces of constant  $W$  therefore move along any path of the particle with velocity  $u = E/mv$ , where  $v$  is the velocity of the particle. This suggests that the velocity  $E/mv$  should be regarded as the wave velocity of the waves composing the particle, and that the surfaces of constant  $W$  are surfaces of constant phase in these waves, for in a train of waves surfaces of constant phase angle move along with the wave velocity. If then  $\psi$  denotes the displacement in these supposed waves we may assume that

$$\begin{aligned} \psi &= a \cos 2\pi\nu \left( \frac{s}{u} - t \right) \\ &= a \cos 2\pi \frac{W}{K} = a \cos \frac{2\pi}{K} (S - Et), \end{aligned}$$

where  $K$  is a constant which has to be introduced since the phase angle must be of zero dimensions, and  $\nu$  is the frequency of vibration. Hence we must have  $E = \nu K$ , and  $S = \nu K s / u$ .

Since in the quantum theory the energy of a quantum is equal to  $h\nu$ , it is natural to assume that  $K = h$ , so that  $E = h\nu$ .

If  $\lambda$  denotes the wave-length then we have  $u = \nu\lambda$ , which with the equations  $u = E/mv$  and  $E = h\nu$  gives  $\lambda = h/mv$ . In the case of a particle describing a circular orbit of radius  $r$  this gives  $\frac{\lambda}{r} = \frac{h}{mvr}$ .

Now in the quantum theory of atoms  $h$  and  $mvr$ , the angular momentum, are of the same order of magnitude, so that it appears that  $\lambda$  and  $r$  are of the same order also. Thus we see that in atomic dynamics we are dealing with phenomena the scale of which is not large compared

with the wave-length, so that the geometrical theory of rays cannot be used but must be replaced by a wave theory.

The equation  $u = \frac{E}{mv}$ , with  $E = h\nu$  and  $v = \sqrt{\frac{2T}{m}} = \sqrt{\frac{2(E-V)}{m}}$ , gives

$$u = \frac{h\nu}{\sqrt{2m(h\nu - V)}}.$$

Thus  $u$  depends on the frequency, so that we have dispersion.

The reciprocal of the group velocity of the waves is  $\frac{d(\nu/u)}{d\nu}$  and so is

$$\frac{d}{d\nu} \left( \frac{\sqrt{2m(h\nu - V)}}{h} \right) = \frac{1}{2h} \frac{2mh}{\sqrt{2m(h\nu - V)}} = \sqrt{\frac{m}{2T}},$$

since  $h\nu - V = E - V = T$ . But  $v = \sqrt{\frac{2T}{m}}$ , so that  $v$  is the group velocity corresponding to the wave velocity  $u$ . Thus we may regard a particle as a group of waves with wave or phase velocity  $u = \frac{E}{mv}$  moving along with velocity  $v$ .

We may suppose that the wave displacement  $\psi$  is determined by the usual wave equation

$$\frac{\partial^2 \psi}{\partial x^2} + \frac{\partial^2 \psi}{\partial y^2} + \frac{\partial^2 \psi}{\partial z^2} - \frac{1}{u^2} \frac{\partial^2 \psi}{\partial t^2} = 0.$$

If we assume  $\psi = \psi_0 e^{2\pi i \nu t}$  we get  $\frac{\partial \psi}{\partial t} = 2\pi i \nu \psi$  and  $\frac{\partial^2 \psi}{\partial t^2} = -4\pi^2 \nu^2 \psi$ , so that

$$\Delta \psi + \frac{4\pi^2 \nu^2 \psi}{u^2} = 0.$$

Now

$$\frac{\nu}{u} = \frac{1}{h} \sqrt{2m(h\nu - V)},$$

so that

$$\Delta \psi + \frac{8\pi^2 m}{h^2} (h\nu - V) \psi = 0.$$

In an atom the wave-length  $\lambda$  is not small compared with the linear dimensions of the atom, so that a particle or electron cannot be regarded as a small body describing orbits inside the atom but must be regarded as a system of waves occupying the whole volume of the atom. In a hydrogen atom the potential energy is  $-e^2/r$  where  $r$  is the distance of the electron from the nucleus. If the electron is a system of waves or vibrations spread over the space around the nucleus it is difficult to see what the potential energy should be. Schrödinger supposes that it is nevertheless equal to  $-e^2/r$  where  $r$  is

the distance of a point from the nucleus. He therefore takes

$$\Delta\psi + \frac{8\pi^2m}{\hbar^2} \left( h\nu + \frac{e^2}{r} \right) \psi = 0$$

to be the differential equation for a hydrogen atom.

Schrödinger succeeded in finding the solutions of this equation, and found that the only solutions for which  $\psi$  is finite and continuous everywhere from  $r = 0$  to  $r = \infty$  give only the following frequencies:

$$\begin{aligned} \nu &> 0, \\ \nu &= -2\pi^2me^4/\hbar^3n^2 \quad (n = 1, 2, 3, 4, \&c.). \end{aligned}$$

The negative sign depends on the negative value taken for the potential energy. This negative value of course is obtained by taking the potential energy zero when  $r = \infty$ , which happens to be convenient for purposes of calculation. The frequencies so calculated are the natural periods of vibration of the atom vibrating like an elastic solid.

The problem of finding the possible frequencies allowed by the equation

$$\Delta\psi + \frac{8\pi^2m}{\hbar^2} \left( h\nu + \frac{e^2}{r} \right) \psi = 0$$

is similar to the problem of finding the possible frequencies of a stretched string fixed at both ends.

If one end of the string is at  $x = 0$  and the other at  $x = l$ , and if  $y$  denotes the lateral displacement of the string, then

$$\frac{d^2y}{dx^2} - \frac{1}{v^2} \frac{d^2y}{dt^2} = 0,$$

where the velocity  $v$  of waves along the string is equal to  $\sqrt{T/\mu}$ ,  $T$  being the tension and  $\mu$  the mass of the string per unit length. The solution of this problem is well known and gives the possible frequencies of vibration of the string or its natural periods. These are found to be given by  $\nu = vn/2l$  where  $n = 1, 2, 3, 4, \dots$ . The fundamental frequency is  $v/2l$ , and the others are multiples of this. In the case of the hydrogen atom the fundamental frequency is  $-2\pi^2me^4/\hbar^3$ , and the others are got by dividing this by 4, 9, 16, &c. The vibrations of the atom, mathematically, extend to  $r = \infty$ , but the amplitude falls off exponentially and becomes negligible when  $r$  is greater than about  $10^{-7}$  cm.

If we put  $\psi = A\epsilon^{-ar}$  and substitute in the differential equation, assuming  $A$  and  $a$  to be constants, we find

$$\Delta\psi = Aa^2\epsilon^{-ar} - \frac{2Aa}{r}\epsilon^{-ar},$$

so that

$$A\epsilon^{-ar} \left( a^2 - \frac{2a}{r} + \frac{8\pi^2m}{\hbar^2} \left( h\nu + \frac{e^2}{r} \right) \right) = 0.$$

Hence 
$$\alpha^2 = -\frac{8\pi^2 m \nu}{h} \quad \text{and} \quad \alpha = \frac{4\pi^2 m e^2}{h^2},$$

so that 
$$\nu = -\frac{2\pi^2 m e^4}{h^3}.$$

Thus in the case of the fundamental vibration of the atom

$$\psi = A e^{\frac{-4\pi^2 m e^2}{h^2} r + 2\pi i \nu t},$$

where  $A$  may have any value and  $\nu = -2\pi^2 m e^4 / h^3$ . The amplitude of vibration at  $r = 0$  is equal to  $A$ . The energy of the atom  $E = h\nu = -2\pi^2 m e^4 / h^2 n^2$  and is independent of the amplitude of vibration. These values of  $E$  are the same as those of Bohr's theory.

According to Schrödinger, the atom radiates when it is vibrating with two of its possible frequencies. For example, if

$$\psi = A_1 e^{2\pi i \nu_1 t} + A_2 e^{2\pi i \nu_2 t}$$

and 
$$\bar{\psi} = A_1 e^{-2\pi i \nu_1 t} + A_2 e^{-2\pi i \nu_2 t},$$

then 
$$\psi \bar{\psi} = A_1^2 + A_2^2 + A_1 A_2 (\epsilon^{2\pi i (\nu_2 - \nu_1) t} + \epsilon^{-2\pi i (\nu_2 - \nu_1) t}).$$

Schrödinger supposes that the density of the electricity  $\rho$  in the atom is proportional to  $\psi \bar{\psi}$ , and that the variations of  $\rho$  produce radiation in accordance with the classical electromagnetic theory.  $\psi \bar{\psi}$  oscillates with frequency  $\nu_2 - \nu_1$ , so that radiation of this frequency is emitted. If then we imagine an atom oscillating with a fundamental frequency  $\nu_2$  and so having energy  $h\nu_2$ , and that it somehow changes from this state to another with frequency  $\nu_1$  and energy  $h\nu_1$ , then if during the transition it has both frequencies it will radiate with frequency  $\nu$  given by  $\nu = \nu_2 - \nu_1$ , and the energy emitted will be

$$h\nu = h(\nu_2 - \nu_1) = E_2 - E_1,$$

which is the fundamental equation of the Bohr theory.

Schrödinger and others have shown that his theory agrees with the facts of spectroscopy as well as or better than Bohr's theory, and it does not require the aid of any correspondence principle. So far as the writer is aware the photo-electric effect has not yet been explained on Schrödinger's new theory.

This undulatory form of dynamics involves a new view of the nature of energy. It seems that the universe consists of a collection of oscillators, the atoms and the light quanta. The energy of any oscillator is equal to  $h\nu$ , so that the principle of the conservation of energy merely means that  $\Sigma \nu$  is constant. The amplitude of vibration of the oscillators has nothing to do with the energy.

A crude analogy is the case of a large collection of tuning forks.

An investigator measures all their frequencies at different times and places, and finds the sum of all the frequencies constant. He then announces the general principle of the conservation of frequency and ignores the amplitude of vibration of the forks.

Schrödinger's theory seems somewhat obscure as yet in parts, but it is undoubtedly a step in the right direction and of the highest interest and importance.

The equation  $\int pdq = nh$  of Bohr's theory may be easily interpreted by means of Schrödinger's theory. If  $s$  is the length of an electron orbit, and  $v$  the velocity along the orbit, we may put  $p = mv$  and  $dq = ds$ , so that  $\int mvd s = nh$ . With the equation  $u = h\nu/mv$  this gives

$$\int \frac{h\nu ds}{u} = nh.$$

But if  $\lambda$  is the wave-length of the waves in the electron,  $\nu\lambda = u$ , so that

$$\int \frac{ds}{\lambda} = n.$$

Now  $\int ds/\lambda$  is the number of waves in the length of the orbit, so that the equation  $\int pdq = nh$  means that the number of waves in the length of the orbit must be equal to  $n$ , which is an integer 1, 2, 3, 4, &c.

On Schrödinger's theory the waves are not regarded as travelling along the orbit only but as forming a system of waves filling up the surrounding space. If  $\int ds/\lambda$  is not equal to an integer, then if the train of waves extends a number of times round the orbit the waves will interfere and destroy each other. In this way we see that the equation  $\int pdq = nh$  only applies to very small orbits the length of which is less than that of the train of waves in an electron.

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## CHAPTER VI

# The Critical Potentials of Atoms

### 1. Quantum Theory and Critical Potentials.

According to the quantum theory atoms can only exist in a series of states characterized by definite energies  $W_1, W_2, W_3 \dots W_n \dots$ . In these so-called stationary states the atom does not emit radiation. The normal state of the atom is the state of smallest energy  $W_1$ . To change an atom from its normal state to another state requires energy  $W_n - W_1$  to be given to the atom. An atom not in the normal state is usually called an excited atom. Excited atoms emit radiation of frequency  $\nu$  given by  $h\nu = W_m - W_n$  when they change from a state having energy  $W_m$  to one having energy  $W_n$ , and they usually quickly revert to the normal state.

If a stream of electrons all moving with the same velocity  $v$  is passed into a gas it is found that the gas atoms may be excited by collisions with the electrons, provided the kinetic energy of the electrons is great enough. The kinetic energy  $\frac{1}{2}mv^2$  must be at least equal to  $W_n - W_1$  for the electrons to be able to change an atom from its normal state with energy  $W_1$  to the state having energy  $W_n$ .

If  $P$  denotes the potential difference required to give an electron the kinetic energy  $\frac{1}{2}mv^2$  so that  $Pe = \frac{1}{2}mv^2$ , where  $e$  is the electronic charge, then if  $Pe = W_n - W_1$  the potential difference  $P$  is called a *critical potential* of the atom for which  $W_m$  and  $W_n$  are possible energies. The potential difference required to give an electron just enough energy to ionize the atom, that is to knock an electron right out of it, is called an ionization potential of the atom. If  $W'$  is the energy of the ionized atom, and  $W_1$  its normal energy, then  $Pe = W' - W_1$  gives the ionization potential of the atom when in its normal state.

### 2. Lenard's Measurements.

The critical potentials of atoms were first directly measured by Lenard with an apparatus shown diagrammatically in fig. 1. Electrons from a hot filament F are attracted by a wire grating or grid G which is charged positively by connecting it to a battery, and some of them pass through the grating. A plate P on the other side of G is connected

to one pair of the quadrants of a quadrant electrometer E, and the other pair of quadrants is connected to the earth. The plate P can be insulated when desired by opening the key K. If  $P_g$  denotes the potential of the grid and  $P_f$  that of the filament, then  $P_g - P_f = V_a$  is called the accelerating potential, and the kinetic energy of the electrons from the filament, when they reach the grid, is approximately given by  $\frac{1}{2}mv^2 = V_a e$ . The potential difference between the grid and

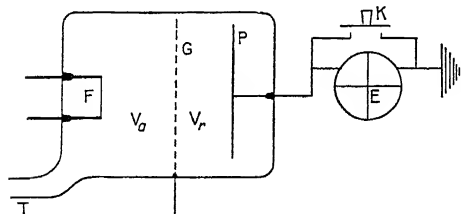


Fig. 1

the plate P is called the retarding potential  $V_r = P_g - P_f$ . The filament is kept at a positive potential less than  $P_g$  so that  $V_r$  is greater than  $V_a$ . The filament, grid, and plate are inside a bulb connected to a pump through a tube T, and the gas

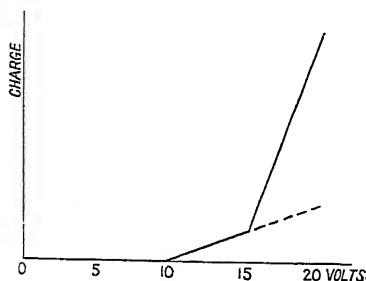


Fig. 2

pressure in the bulb is adjusted so that the mean free path of the electrons is greater than the distance from the filament to the grid. The electrons which pass through the grid are repelled by the plate and, since  $V_r > V_a$ , none of them can get to it. If, however, the energy of the electrons is great enough they ionize some of the gas molecules by collisions in the space between the grid and plate, and the positive ions produced are attracted by the plate P and give it a positive charge which is indicated by the electrometer when the key

K is open. If  $V_a$  is gradually increased, keeping  $V_r$  always greater than  $V_a$ , then the smallest value of  $V_a$  for which the plate gets any positive charge can be found.

Fig. 2 shows the relation between  $V_a$  and the positive charge received by the plate in a given time with hydrogen gas in the apparatus. No charge is observed until  $V_a$  is about 10 volts. The charge increases slowly up to about 16 volts and then begins to increase rapidly with the accelerating potential. This shows that hydrogen has critical potentials of 10 and 16 volts approximately.

The positive charge received by the plate P, however, is not necessarily due to positive ions; it may be caused by ultra-violet light emitted by the gas falling on the plate and causing it to emit electrons. The electrons from the filament may have enough energy to excite the gas molecules but not enough to ionize them. If the molecules

are excited by the electron collisions they emit light when they go back to the normal state, and this light may cause the plate to emit electrons and so become positively charged. Lenard's method shows that the hydrogen has critical potentials at about 10 and 16 volts, but it does not enable us to determine whether these potentials are ionization potentials or excitation potentials.

### 3. Results of Franck and Hertz with Mercury Vapour.

A great many modifications of Lenard's original method have been devised but only a few of these can be described here. A full account is given in the *Bulletin of the National Research Council*, No. 48, "Critical Potentials", by K. T. Compton and F. L. Mohler.

If the gas pressure in the apparatus is increased so that the mean free path of the electrons becomes considerably smaller than the distance from the filament to the grid and if also the retarding potential is made quite small (about  $\frac{1}{2}$  volt), then, with monatomic gases, the current to the plate rises and falls periodically as the accelerating potential is increased. Results obtained in this way by Franck and Hertz with mercury vapour are shown in fig. 3.

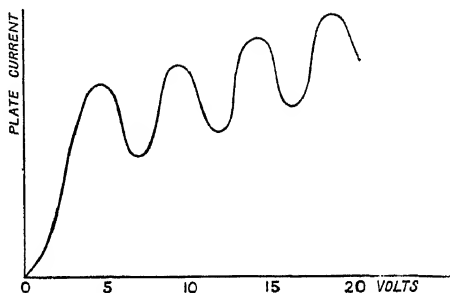


Fig. 3

The potential differences between the successive maxima are 4.9 volts. It is supposed that the electrons, when their kinetic energy corresponds to less than 4.9 volts, collide with the mercury atoms without loss of energy. The collisions are then said to be elastic. An electron from the filament therefore loses no energy due to collisions as it moves towards the grid until its energy corresponds to 4.9 volts, and then it excites a mercury atom and loses all its kinetic energy. The energy of the electron then increases again as it moves along in the electric field until it again corresponds to 4.9 volts, when it again excites a mercury atom and loses its energy, and so on. In order to get to the plate an electron must arrive at the grid with energy greater than that corresponding to the small retarding potential  $V_r$ . Hence as  $V_a$  is gradually increased the current increases until  $V_a$  is about 4.9 volts, when it drops, because the electrons lose their energy just before reaching the grid. The current then increases again as  $V_a$  is increased until it reaches  $2 \times 4.9$  volts, when it again falls and so on. The lowest critical potential of mercury atoms is therefore 4.9 volts. Similar results have been obtained with helium and other monatomic gases.

More precise results can be obtained by using two grids, one near the filament and the other near the plate. The one near the plate is kept at a slightly higher potential (about 0.1 volt) than the one near the filament. The electrons which pass through the first grid then move with nearly constant velocity in the space between the two grids, so that the number of collisions occurring with any given energy is greatly increased, and the breaks in the curve giving the relation between plate current and accelerating potential are much more distinct. The second grid can be used with Lenard's method at low pressures and with Franck and Hertz's method at higher pressures. In this way a great many critical potentials

have been observed; for example, in mercury there are critical potentials of 4.68, 4.9, 5.29, 5.78, and 6.73 volts.

#### 4. Methods distinguishing Ionization and Excitation Potentials.

A modification which enables ionization and excitation potentials to be easily distinguished was used by K. T. Compton and by Boucher. Two plates were provided, one consisting of a metal sheet, as usual, and the other of a grating of fine wire. The two plates were connected together and either of them could be moved into position in front of the grids when desired. The area of the metal sheet was much greater than that of the grating so that the current due to radiation was much greater with the sheet than with the grating, but the current due to positive ions was about the same with either plate. The ratio of the plate current with the sheet to the plate current with the grating remained nearly constant as the accelerating potential was increased

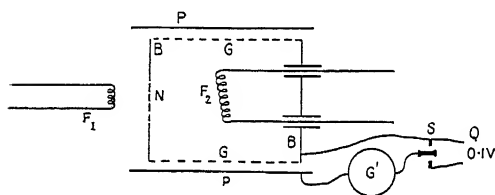


Fig. 4

so long as only excitation of atoms was taking place, but began to fall rapidly as soon as ionization began.

Fig. 4 shows diagrammatically an apparatus due to Hertz which enables two dif-

ferent methods to be used. BB is a cylindrical box with flat ends. In one end at N is a hole covered with a grating. The sides of the box GG are made of wire gauze. Surrounding the box is a metal cylinder PP. A filament  $F_1$  is set up near the hole at N and there is a second filament  $F_2$  inside the box.

To determine ionization potentials the filament  $F_2$  is heated and the current from box to filament due to a small potential difference is measured. The temperature of  $F_2$  is raised until the current is independent of the temperature, showing that the current is limited by the space charge on the electrons emitted. If now electrons from the other filament  $F_1$  are accelerated into the box through the hole at N by a potential difference  $V_a$ , they have no effect on the current unless they produce positive ions in the box by collisions with the gas molecules present. The positive ions neutralize some of the space charge and so increase the current. Owing to the mass of the positive ions being much greater than that of the electrons the ions remain in the space inside the box much longer than the electrons, so that the increase of current is enormously greater than the current carried by the positive ions. This is therefore a sensitive method for detecting an ionization potential.

The same apparatus can also be used to detect any kind of critical

potential by another method. The filament  $F_2$  inside the box is not used but is merely connected to the box. Electrons from  $F_1$  are accelerated into the box through N and some of them escape from the box through the gauze sides to the cylinder P. The current to P carried by these electrons is measured with a galvanometer G' connected to P. A switch S is provided in the galvanometer circuit by means of which a potentiometer Q arranged to give a small P.D. of about 0.1 volt can be introduced into the circuit when desired. The current is observed with the small P.D. on, and also with it off. With the P.D. on, the electrons from the box cannot get to P unless they have enough energy to carry them across the small P.D., but with the P.D. off all the electrons from the box get to P. The difference between the galvanometer currents with the P.D. off and with it on therefore gives the number of electrons coming out of the box which have kinetic energies between zero and the energy corresponding to the small P.D.

If  $V_a$  is the accelerating potential driving the electrons from  $F_1$  into the box and if  $V_c$  is a critical potential for the gas in the box, then if  $V_a - V_c$  is between zero and the value of the small P.D. there will

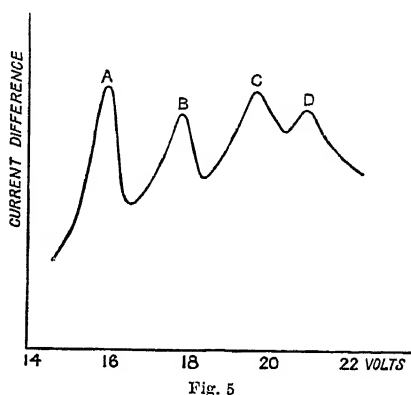


Fig. 5

be electrons in the box with energies between zero and that corresponding to the small P.D., some of which will escape from the box and produce a difference between the two currents observed. If, however,  $V_a - V_c$  is greater than the small P.D. or less than zero there will be no such electrons present and the two currents will be equal.

As the accelerating potential is gradually increased the two currents remain equal until a critical potential is reached, when a difference appears which rises to a maximum and falls back to zero when  $V_a - V_c$  becomes greater than the small P.D. On  $V_a$  being further increased, when a second critical potential is reached a difference between the two currents appears again and then disappears, and so on. Fig. 5 shows the results obtained by Hertz with a mixture of neon and helium. The two maxima at A and B are due to critical potentials of helium and those at C and D to neon, as can be shown by trying the two gases separately.

All the above methods are subject to an error on account of the initial velocity of the electrons emitted by the hot filament. The energy of the electrons is greater than that corresponding to the

accelerating potential  $V_a$ . This error is best determined by measuring known critical potentials, or by measuring the difference between the critical potential which it is desired to find and a known critical potential. Possible errors due to contact potential differences are also eliminated in this way.

### 5. Agreement of Results with Quantum Theory.

By means of the above and other more or less similar methods the critical potentials of many different atoms and molecules have been determined. These results provide a remarkable confirmation of one of the fundamental assumptions of the quantum theory, viz. that the energy of atoms does not vary continuously but that atoms can only exist in a series of definite states having definite energies. Electrons collide elastically with an atom unless they have enough kinetic energy to change the energy of the atom from its actual state to another possible state.

According to Bohr's quantum theory of spectra, moreover, we have

$$h\nu = W_m - W_n,$$

so that if  $P$  denotes the critical potential corresponding to the change from energy  $W_n$  to energy  $W_m$ , we have

$$h\nu = Pe.$$

If  $P$  is expressed in volts and  $\nu$  as the wave number or number of waves in 1 cm. in a vacuum, then on substituting the values of  $h$  and  $e$  we get

$$P = 1.2344 \times 10^{-4} \nu.$$

It is found that the observed critical potentials in many cases agree accurately with the values calculated by means of this equation. Since the wave number can be determined with much greater accuracy than the critical potential  $P$ , it follows that when the spectral line corresponding to a particular critical potential has been definitely identified, then the value of the critical potential can be most accurately obtained by calculation from the wave number of the spectral line.

The results which have been obtained with the alkali metals are especially simple. The vapours of these metals are monatomic, and are found to absorb the lines of the principal series in their spectra strongly. This shows that the lines of this series are emitted when the atoms revert to the normal state from a series of states having greater energies. See Chap. VIII.

An excitation potential has been found for these atoms which agrees with that calculated from the wave number of the first line in the principal series, and also an ionization potential which agrees with that calculated from the wave number of the limit of the principal

series. The following table gives the observed and calculated results in volts.

Metal.	Wave Number of Limit.	Ionization Potential.		Wave Numbers of First Line.	Excitation Potential.	
		Cal.	Obs.		Cal.	Obs.
Sodium ..	41,449	5.116	5.13	{ 16,973 16,956	{ 2.095 2.093	2.12
Potassium	35,006	4.321	4.1	{ 13,043 12,985	{ 1.610 1.603	
Rubidium	33,689	4.159	4.1	{ 12,817 12,579	{ 1.582 1.553	1.6
Cæsium ..	31,405	3.877	3.9	{ 11,732 11,178	{ 1.448 1.380	

The first lines of the principal series are doublets, but the measurements of the excitation potentials were not sufficiently precise to distinguish between the two critical potentials.

Excitation potentials corresponding to the other lines in the principal series have not yet been observed. We should expect the ionization potential to agree with that calculated from the limit of the principal series because the successive lines are supposed to correspond to states in which the outer electron is farther and farther from the nucleus, so that the limit should correspond to the case when the electron is entirely removed from the atom.

The following table contains the observed and calculated critical potentials of helium in volts.

Calculated.	Observed.
19.73	.... 19.73
20.56	.... 20.53
21.12	.... 21.2
22.98	.... 22.9
24.48	.... 24.5

The calculated values are got from the wave numbers of spectral lines believed to be emitted when the helium atom reverts to its normal state, except the last number, which is got from the limit of the corresponding series, since 24.5 volts is the ionization potential.

In the case of mercury, eighteen critical potentials have been observed ranging from 4.68 volts to the ionization potential 10.39 volts. Most of these agree with values calculated from the wave numbers of spectral lines believed to be emitted by mercury atoms reverting to the normal state. Similar results have been obtained with many other elements.

## 6. Smyth's Application of Positive Ray Analysis.

Very interesting and important results have been obtained by Smyth by means of a combination of Lenard's method of measuring critical potentials and positive ray analysis. In this way it is possible to determine the nature of the positive ions produced in gases by electron collisions.

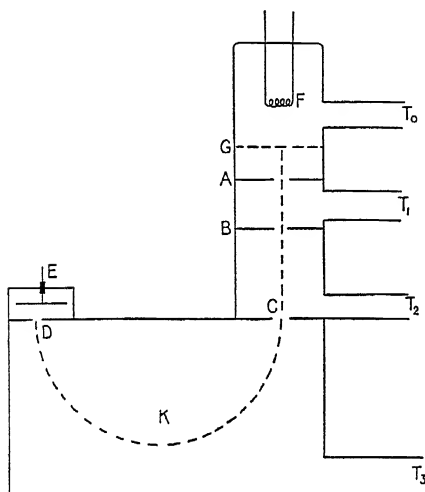


Fig. 6

insulated electrode E. The charge received by E is measured with a quadrant electrometer. The box K and the spaces between the diaphragms A, B, and C are kept highly exhausted by means of powerful pumps connected to the tubes  $T_1$ ,  $T_2$ , and  $T_3$ , so that the positive ions are not stopped by collisions with gas molecules.

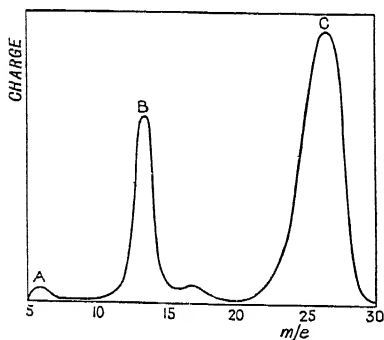


Fig. 7

A slow stream of the gas to be investigated is passed into the apparatus through the tube  $T_0$ , making it possible to have sufficient gas pressure in the space between the grid and plate to obtain enough ions by collisions although the rest of the apparatus is kept highly exhausted.

The accelerating potential  $V_a$  between the filament and grid is kept constant, as also the retarding potential between G and A. The potential between A and B which accelerates the positive ions produced between G and A is varied and the magnetic field kept constant. The values of  $m/e$  for the positive ions can be easily calculated from the radius of the path which they describe, the potential accelerating them, and the strength of the magnetic field.

In fig. 7 the way in which the charge received by the electrode E varies with the value of  $m/e$  for the positive ions reaching E is shown. In this case the gas used was nitrogen and the accelerating potential  $V_a$  was 100 volts. The value

of  $m/e$  is expressed in terms of  $m/e$  for protons, so that it is equal to the atomic weight in the case of ions carrying a single electronic charge.

The curve shows three maxima at  $m/e$  equal to 7, 14, and 28. These must clearly be due to positive ions consisting of atoms of nitrogen with two electronic charges, atoms with one electronic charge, and molecules  $N_2$  with one electronic charge. The relative amounts of these three sorts of positive ions given by nitrogen was determined for a series of values of the accelerating potential  $V_a$ . It was found that when  $V_a$  is less than 25 volts only molecules  $N_2^+$  with one electronic charge are obtained. Above 25 volts small amounts of  $N^+$  and  $N^{++}$  appear. This shows that when nitrogen gas is ionized by collisions with electrons practically all the positive ions produced are  $N_2^+$ . The nitrogen molecule is evidently a very stable system not easily dissociated into two atoms. This result explains the chemical inertness of nitrogen.

Very similar results were obtained with hydrogen.  $H_2^+$  ions were readily formed but  $H^+$  was obtained more easily than  $N^+$ .

The critical potentials of atoms corresponding to the production of characteristic X-rays are discussed in the chapter on X-rays.

#### REFERENCE

"Critical Potentials." K. T. Compton and F. L. Mohler. *Bulletin of the National Research Council, Washington, D.C.*

## CHAPTER VII

# X-Rays and $\gamma$ -Rays

### 1. Nature of X-rays.

X-rays or Röntgen rays were discovered by Röntgen in 1895. He was using a Crookes tube in which the cathode rays struck the glass walls of the tube, and he noticed that a piece of paper coated with barium-platinocyanide, which happened to be lying near, fluoresced when the tube was working. On putting objects between the tube and the paper, shadows were obtained in the fluorescent light, showing that the tube was emitting some kind of radiation. It was found that this radiation was remarkably penetrating. It passed readily through black paper and thin sheets of aluminium which were quite opaque to ordinary light. It was found that the new radiation could not be refracted or reflected but travelled along straight lines through objects of any shape, and it was not deflected by either magnetic or electric fields.

J. J. Thomson discovered that the X-rays produce electrical conductivity in gases and other insulators when passed through them.

It was found that the X-rays are emitted where the cathode rays strike any solid object. In 1881 J. J. Thomson pointed out that if cathode rays are rapidly moving charged particles, then, according to Maxwell's electromagnetic theory, they should produce electromagnetic radiation when suddenly stopped. He suggested that the green fluorescence of the glass walls of Crookes tubes produced by the cathode rays may be produced by this radiation emitted when the rays strike the glass. Soon after the discovery of X-rays Stokes and Wiechert suggested that the sudden stopping of the cathode rays when they strike a solid body causes the emission of electromagnetic pulses or waves of very short wave-length which are the X-rays. It is believed now that the cathode rays penetrate the atoms of the solid body and are not stopped suddenly. The cathode rays ionize and excite the atoms by collisions and the X-rays are emitted when the atoms revert to their normal state. X-rays are simply light or electromagnetic waves of very short wave-length. Barkla in 1905 showed that polarized X-rays can be obtained, and it is now known that X-rays can be

diffracted, reflected, and refracted, and so have all the properties of light on which the wave theory of light is based.

## 2. The Coolidge Tube.

The X-rays from ordinary Crookes tubes are very feeble, and powerful X-rays are now obtained from specially designed tubes. The most satisfactory X-ray tubes are the Coolidge tubes, in which the electrons emitted by a hot filament are accelerated towards a metal anode usually made of tungsten or molybdenum.

A Coolidge tube is shown diagrammatically in fig. 1.

The cathode C consists of a small flat spiral of tungsten wire surrounded by a tungsten cylinder. The spiral is heated by a current

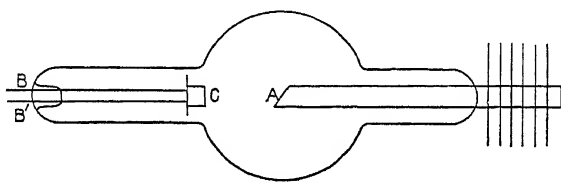


Fig 1

from a battery through the wires B and B' sealed through the glass. The anode A consists of a plug of tungsten or molybdenum set in the end of a copper rod which is cut off at 45° as shown. The other end of the copper rod, outside the bulb, is cooled by means of several copper discs which expose a large surface to the air. The bulb is very completely exhausted, so that no electrical discharge can be passed through it when the filament is cold.

A difference of potential of about 50,000 to 100,000 volts is maintained between the anode and cathode, and the filament heated so that it emits electrons. The current through the tube can be measured with a milliammeter and may be up to 50 or 100 milliamperes. The electrons emitted by C are concentrated on to a small area on the end of A by the electrostatic repulsion of the charge on the end of the cylinder surrounding the filament. The X-rays are emitted by this small area about equally in all directions. They are absorbed of course by the anode when they go into it. The anode of a tube taking, say, 50 milliamperes at 100,000 volts or 5 kilowatts rapidly gets hot, for nearly all the energy of the electrons is converted into heat when the electrons are absorbed by the anode. Less than 1 per cent of the energy goes into the X-rays.

## 3. Scattering. Diffraction. Characteristic X-rays.

When X-rays are passed through matter they are partly scattered in all directions, while at the same time the matter may be caused to

emit secondary X-rays different in quality from the incident rays. Barkla discovered that the secondary X-rays emitted by different elements have properties characteristic of the particular element emitting them which vary in a regular way with the atomic weight of the element. X-rays also cause matter to emit electrons, an effect analogous to the emission of electrons due to ultra-violet light.

In 1912 Laue discovered that X-rays can be diffracted by crystalline substances, which act on the rays in the same sort of way as a diffraction grating acts on light. By means of this effect it is possible to determine the wave-lengths of X-rays, and it is found that the X-rays emitted by different elements have definite wave-lengths characteristic of the element, just like the wave-lengths of the spectral lines in the optical spectra of the elements. Laue's important discovery placed the study of X-rays on a new quantitative basis and will now be considered in detail.

## X-RAYS AND CRYSTAL STRUCTURE

### 4. Crystallography. Law of Rational Indices.

Crystals are homogeneous solid bodies bounded by plane surfaces which are called the faces of the crystal. The faces are arranged in sets, all those in a set being perpendicular to the same plane; these sets are called zones. The faces in a zone intersect in parallel lines. The faces occur in parallel pairs on opposite sides of the crystal.

The geometrical character of the crystals of any substance is determined by the angles between their faces, which are invariable, and is independent of the size of the particular crystal selected. Crystals grow by the deposition of uniform layers on their faces, so that they get larger without the angles between the faces changing.

The different faces of a crystal can be specified by the intercepts cut off from three axes by the faces. As axes, lines parallel to three edges of the crystal which do not lie in one plane are used.

In fig. 2, let OA, OB, and OC be the three axes selected, and let E, F, G be the points at which a face of the crystal intersects them. Let OE =  $a$ , OF =  $b$ , and OG =  $c$ . The ratios  $a : b : c$  will always be the same for the corresponding axes and face on any crystal of the same substance, whatever the size of the crystal. Any other plane parallel to the face EFG will cut off lengths from the axes in the same ratios as  $a : b : c$  and could equally well be taken to be the face of the crystal.

Now let  $a'$ ,  $b'$ , and  $c'$  be the intercepts for any other face of the crystal; then it is found that there is always a simple numerical relation between  $a'$ ,  $b'$ ,  $c'$  and  $a$ ,  $b$ ,  $c$ . The ratios  $a' : b' : c'$  are equal to the ratios  $a/k : b/h : c/l$ , where  $k$ ,  $h$ , and  $l$  are integers which are usually quite small and are called the indices of the face they specify. Thus

$$\frac{a/k}{b/h} = \frac{a'}{b'}, \text{ so that } \frac{h}{k} = \frac{b}{a} \frac{a'}{b'}$$

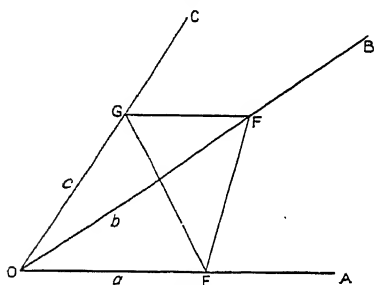


Fig. 2

and in the same way

$$\frac{l}{h} = \frac{c}{b} \frac{b'}{c'} \text{ and } \frac{k}{l} = \frac{a}{c} \frac{c'}{a'}.$$

If  $k = h = l = 1$ , we have  $a' : b' : c' = a : b : c$ , and so have the original face. If  $k = h = 1$  and  $l = 2$  we have  $a' : b' : c' = a : b : c/2$ ; and if  $k = 1, h = 2$ , and  $l = 3$  we have  $a' : b' : c' = a : b/2 : c/3$ , and so on. Thus any face of the crystal can be specified by giving the values of  $k, h$ , and  $l$  for that face. The values of  $k, h, l$ , of course, depend on the particular edges chosen as axes and on the particular face selected as the  $k = h = l = 1$  face.

If a face is parallel to one of the axes, as frequently happens, then its intercept on that axis is infinitely long, so that the integer becomes zero. Thus if  $k = 0$ ,  $h = 1, l = 1$  so that  $a' = a/0 = \infty, b' = b, c' = c$ , the face is parallel to the axis OA. The indices  $k, h$ , and  $l$  can be positive or negative integers or zero. Any face of a crystal can therefore be specified by giving the values of  $h, k$ , and  $l$  for that face. The numbers are usually put inside a bracket, thus (112) denotes the face with intercepts  $a, b$ , and  $c/2$ . Negative values are indicated by a minus sign above the number thus ( $\bar{1}\bar{1}\bar{2}$ ).

By properly selecting the axes and reference face (111) it is usually possible to represent all principal faces by quite small values of  $h, k$ , and  $l$ .

As an example, consider a cube (fig. 3) and let the origin be at its centre and the axes parallel to three of its edges which meet at one corner. In this case, if  $a = b = c$ , the six sides of the cube are (100), ( $\bar{1}$ 00), (010), (0 $\bar{1}$ 0), (001), (00 $\bar{1}$ ), since each side is perpendicular to one axis and parallel to the other two. The reference face EFG passes through the three points where the positive directions of the axes cut the surface of the cube. It is not one of the faces of the cube, but is a possible face for a crystal having faces parallel to the sides of the cube. Eight possible faces are (111), ( $\bar{1}\bar{1}$ 1), (1 $\bar{1}\bar{1}$ ), ( $\bar{1}\bar{1}$ 1), ( $\bar{1}\bar{1}$ 1), (1 $\bar{1}\bar{1}$ ), ( $\bar{1}\bar{1}$ 1), ( $\bar{1}\bar{1}$ 1). These eight faces are shown in fig. 4. They form an octahedron.

The fact that the indices are positive or negative integers or zero for any face of a crystal is known as the law of rational indices. This law, which was first discovered empirically, can be explained in a simple way on the atomic theory. We suppose that crystals consist of a regular arrangement of atoms, the same throughout the volume of the crystal. The atoms are arranged in groups or sets, all groups being similar and similarly orientated. In a crystal of a compound

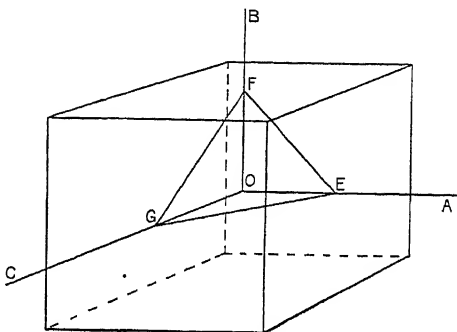


Fig. 3

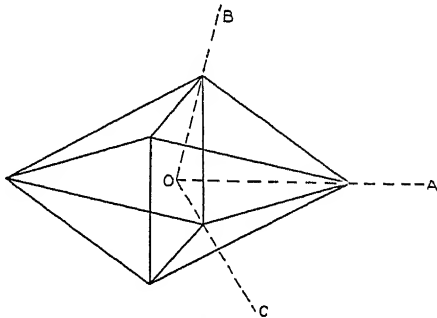


Fig. 4

body each group will usually consist of the atoms forming one molecule of the compound. In crystals of an element each group may consist of a single atom. The position of a group is fixed by the position of any definite point in it, for example, the centre of one of the atoms in it. We shall for the present speak of each group as being located at the point fixing its position. The groups must be arranged in the crystal in such a way that every group is related to its neighbouring groups in the same way. Thus if  $A_3$  and  $A_4$  are two groups in a crystal (fig. 5) then there must also be groups at  $A_1, A_2, A_5, A_6$  on the line joining  $A_3$  and  $A_4$ ,

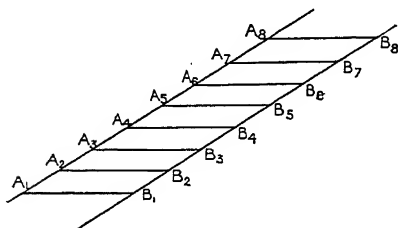


Fig. 5

and the groups must be equally spaced along this line. If  $B_1$  is another group near  $A_1$ , then there must also be groups at  $B_2, B_3, B_4$ , &c., in such positions that  $A_2B_2 = A_1B_1$ ,  $A_1A_2 = B_1B_2$ , &c. In this way it is easy to see that all the groups in a crystal must be arranged at the intersections of three sets of equally spaced parallel planes. Such an arrangement is called a space lattice.

Three sets of equally spaced parallel planes divide space up into equal parallelepipeds as shown in fig. 6, and there is one group of atoms at each point where three planes intersect. The number of groups of atoms is equal to the number of the parallelepipeds.

If a plane is drawn through any three groups of atoms in the space lattice, the plane will contain groups of atoms arranged at the intersections of two sets of parallel lines, and it is supposed that such a plane is a possible face of the crystal. The faces usually observed are those determined by planes which pass through three groups of atoms which are near together in the space lattice. If three sides of one of the parallelepipeds, meeting at a point, are taken as the

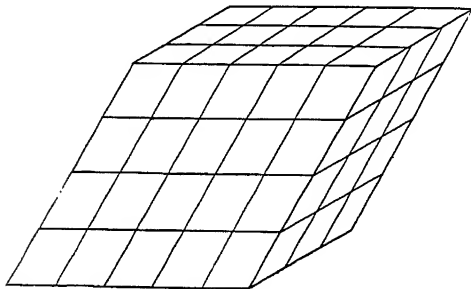


Fig. 6

axes then any plane drawn through three groups will have intercepts on these axes which are multiples of the lengths of the sides of the parallelepipeds. Let these lengths be  $a_1, a_2, a_3$  so that the intercepts are  $n_1a_1, n_2a_2, n_3a_3$ , where  $n_1, n_2, n_3$  are integers. For another plane through three other groups let the intercepts be  $n'_1a_1, n'_2a_2, n'_3a_3$ . If we put  $n_1a_1 : n_2a_2 : n_3a_3 = n'_1a_1/k : n'_2a_2/h : n'_3a_3/l$ , we have  $h/k = n'_2n_1/n'_1n_2$ ,

$l/h = n_3n'_2/n_2n'_3$ ; but the  $n$ 's are all integers and therefore the ratios  $h/k$  and  $l/h$  can be expressed as the ratios of integers. Thus we see that the law of rational indices follows from the theory that crystals consist of groups of atoms arranged in space lattices and that the faces are planes drawn through the groups. The axes and reference face (111) are chosen for any particular crystal so that the other faces are given by the simplest possible sets of indices. The principal faces of a crystal can usually be specified without using integers greater than 2. It should be observed that there are many different sets of equally spaced parallel planes the intersections of which form any particular space lattice.



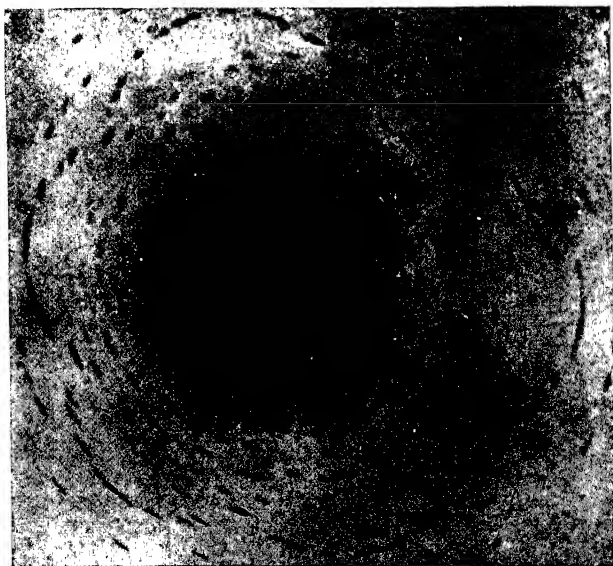


Fig. 8, Chap. VII.—Diffraction of X-Rays by a Crystal of Beryl  
(From *X-Rays and Crystal Structure*, Sir W. H. and W. L. Bragg)

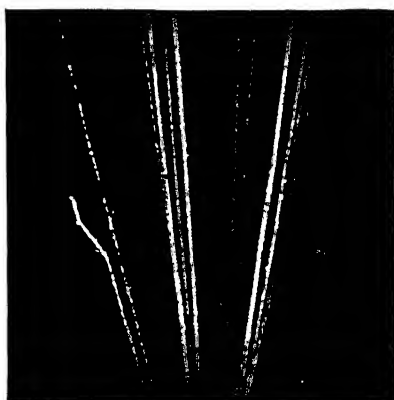
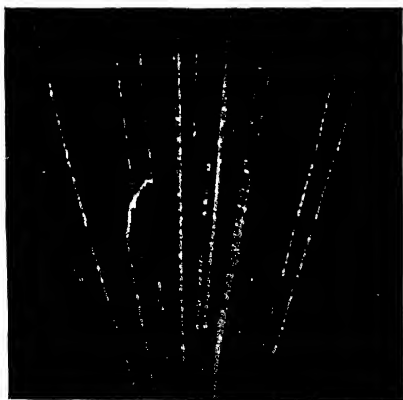


Fig. 5, Chap. XIII.—Photographs of  $\alpha$ -ray tracks through nitrogen, by P. M. S. Blackett, showing collision between  $\alpha$ -ray and nitrogen atom, at which a track branches into a thin straight track due to the ejected proton and a thick track due to the nitrogen atom combined with the  $\alpha$ -ray. See p. 225.

### 5. Diffraction Patterns.

If we draw planes through every group in a crystal, all parallel to one of the faces, we get a set of equally spaced parallel planes. Since all the groups of atoms are supposed to be similar and similarly orientated it follows that the arrangement of atoms is repeated at regular equal intervals measured from the face along any normal to it. When X-rays are passed through matter they are partly scattered in all directions. It is supposed that this scattering is produced by the electrons in the atoms. If the X-rays are a train of waves of definite wave-length  $\lambda$ , each electron is supposed to be made to vibrate with the frequency  $c/\lambda$  of the rays by the electric field in the rays. The electron therefore emits radiation of the same wave-length as the rays falling on it.

If a narrow beam of X-rays is passed through a crystal, then in certain directions the scattered rays from the regularly arranged atoms are all in phase and together form a comparatively intense scattered beam. The apparatus used in Laue's original experiment is shown in fig. 7. The rays from a tube T are passed through a series of small holes in lead screens A, B, C, and the narrow pencil of rays falls on a crystalline plate P about 1 mm. thick. A photographic plate S is put up a few centimetres behind the crystal.

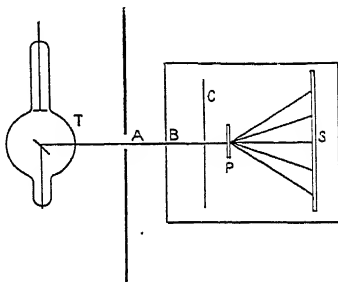


Fig. 7

After an exposure of several hours to the X-rays, on developing the plate a symmetrical pattern of spots is found on it. These spots are arranged round a central spot formed by the rays which pass straight through the crystal. Some of the diffracted rays may make angles of  $40^\circ$  or more with the incident pencil. Fig. 8 shows reproductions of such diffraction patterns. (See Plate.)

### 6. W. L. Bragg's Theory of Diffraction Patterns.

The theory of these diffraction patterns was worked out by Laue and was found to agree exactly with the facts. Laue's theory was complicated, and a greatly simplified form of the theory was proposed by W. L. Bragg. Bragg's theory gives the same results as Laue's and will be considered here.

Consider a narrow beam of X-rays from a source  $S$  (fig. 9) falling on a plane  $EF$ , and suppose that single atoms are distributed over this plane. Each atom scatters a minute fraction of the rays falling on it in all directions. Suppose that the scattered waves from two atoms at  $A$  and at  $A'$  which are near together arrive at a point  $P$  at the same

time so that the waves reinforce each other at  $P$ . For this to happen we must have

$$SA + AP = SA' + A'P.$$

Now if  $SA + AP = \text{constant}$ ,  $A$  must be on the surface of an ellipsoid of revolution about  $SP$  with foci at  $S$  and  $P$ . If the plane  $EF$  is a tangent plane to this ellipsoid at  $A$ , the scattered waves from all the atoms on the plane  $EF$  which are very near  $A$  will arrive at  $P$  together, so that there will be regular reflection of a very small fraction of the X-rays along the path  $SAP$ . If  $EF$  is a tangent plane to the ellipsoid, then  $SA$ ,  $AP$ , and the normal  $AN$  to  $EF$  at  $A$  lie in the same plane, and the angle  $SAN$  is equal to the angle  $NAP$ .

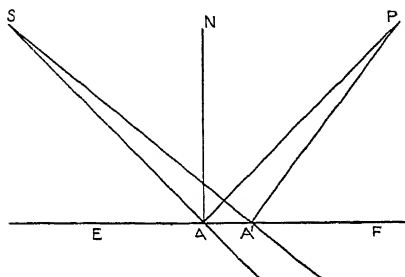


Fig. 9

Now suppose that the plane  $EF$  is on the face of a crystal. As we have seen, the atoms in the crystal are regularly arranged in groups about equally spaced parallel planes of which the face is one. Let the distances between these equally spaced plane layers of atoms be  $d$ . Then each layer will regularly reflect a small fraction of the rays falling on it. If the rays reflected from all the layers are in the same phase and so reinforce each other a strong reflected beam will be obtained. In fig. 10 let  $S$  be the source of the rays and  $A_1, A_2, A_3$ , &c.,

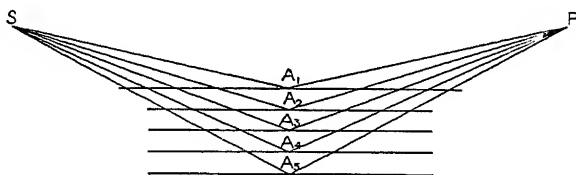


Fig. 10

the equally spaced layers of atoms. Let the rays from  $S$  be reflected at  $A_1, A_2, A_3$ , &c., to  $P$ , and suppose that the reflected wave trains from each plane arrive at  $P$  in the same phase and so reinforce each other at  $P$ . This requires that

$$(SA_2 + A_2P) - (SA_1 + A_1P) = n\lambda,$$

$$(SA_3 + A_3P) - (SA_2 + A_2P) = n\lambda,$$

$$(SA_4 + A_4P) - (SA_3 + A_3P) = n\lambda,$$

. . . . .

where  $n$  is a positive integer; that is, the path differences for the successive planes must be a multiple of the wave-length  $\lambda$ . The distance  $d$  between the reflecting planes is very small compared with  $SA$  and  $AP$  so that the rays near  $A$  are practically parallel. The path differences are therefore easily calculated. In fig. 11 let  $p$  and  $p'$  be two successive reflecting planes so that  $AA' = d$ , and let the angles  $ANA'$  and  $AMA'$  be right angles. The path difference is  $NA' + A'M$ , which is equal to  $2d \sin \theta$  where  $\theta$  is the angle between the incident or the reflected rays and the reflecting planes. The condition for strong reflection from the crystal face is therefore

$$2d \sin \theta = n\lambda,$$

where  $n = 1, 2, 3 \dots$

Such reflection of X-rays at a crystal face, of course, is not a reflection at the surface of the crystal but is reflection by a very large number of the equally spaced layers of atoms. The distances between the layers are of the order of magnitude  $10^{-8}$  cm., so that there are a million layers in a thickness of the order of  $\frac{1}{10}$  mm. The angle  $\theta$  is usually called the glancing angle, and  $d$  is often called the crystal grating space. There is no appreciable reflection at angles differing even very slightly from the values given by  $2d \sin \theta = n\lambda$ . For if the path difference is  $\lambda \left( n + \frac{1}{2m} \right)$ , instead of  $n\lambda$ , where  $m$  is a large whole number, then there is a path difference  $\lambda \left( nm + \frac{1}{2} \right)$  between the rays reflected from any plane and the  $m$ th plane below it. The number  $nm$  is an integer, so that the two wave trains interfere and destroy each other. The reflected wave trains therefore interfere in pairs and there is no appreciable reflection, unless  $m$  is comparable with the total number of reflecting planes, which is very large. When a beam of X-rays is passed through a crystal as in Laue's experiment, the rays may be reflected from all the sets of planes into which the groups of atoms can be supposed arranged. If the X-rays are not homogeneous but have a continuous spectrum so that all wave-lengths between certain limits are present, then each set of planes reflects the rays of wave-length  $\lambda$  given by

$$2d \sin \theta = n\lambda,$$

where  $\theta$  is the glancing angle between the set of planes and the incident beam. In this way the diffraction patterns obtained can be explained.

The intensity of the reflection from any set of layers increases with the number of atoms per square centimetre in the layers, so that, other

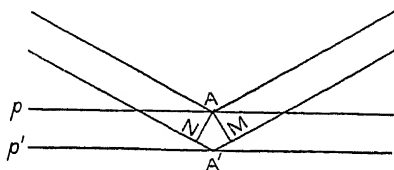


Fig. 11

things being equal, sets for which  $d$  is small reflect less than those for which  $d$  is large. The more intense spots in the patterns are those from sets of planes which can be drawn through three groups of atoms which are close together in the crystal.

### 7. W. H. Bragg's X-ray Spectrometer.

The reflection of X-rays from the faces of crystals enables the wavelengths of the rays to be found and also the arrangement of the atoms in the crystal. This method of studying X-rays and crystals is due to W. H. Bragg. The instrument used for measuring the glancing angles is called an X-ray spectrometer.

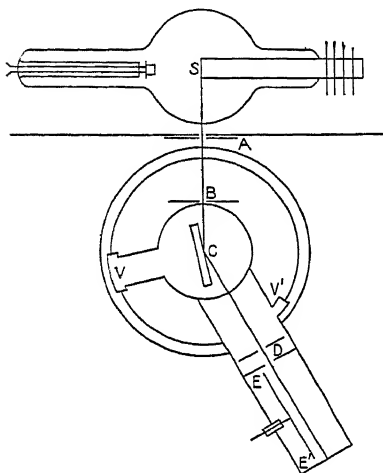


Fig. 12

Fig. 12 shows the essential parts of an X-ray spectrometer used by W. H. Bragg. The X-rays from the anti-cathode S of a Coolidge tube pass through two slits A and B and fall on the face of a crystal at C. The crystal is mounted on a table which can be rotated about an axis in the crystal face and the angles through which it is turned can be read off on a scale by the vernier V. The reflected rays pass through a slit at D and then through a thin aluminium window into an ionization chamber EE'. The slit D and chamber

can be turned about the same axis as the crystal and the angles read off by the vernier V'.

The ionization chamber is a cylindrical lead box about 15 cm. long and 5 cm. in diameter. The narrow beam of rays passes along the centre of the chamber and ionizes the gas in it. The chamber contains an insulated electrode, which is connected to some form of sensitive electrometer. C. T. R. Wilson's tilted gold leaf electroscope is frequently used. The chamber is charged to about 100 volts by connecting it to a battery, and the insulated electrode is kept near zero potential. The conductivity produced in the chamber by the X-rays can be measured with the electrometer. The chamber is usually filled with ethyl bromide vapour or some other gas which absorbs the rays strongly.

To study the reflection of the X-rays by the crystal, the chamber slit is set so as to receive the reflected rays. The angle between the chamber slit and the incident rays must be twice the angle between the

crystal face and the incident rays. If the ionization is measured for a series of glancing angles a curve may be drawn showing the variation of ionization with the angle. Fig. 13 shows such a curve obtained by Bragg with the X-rays from a rhodium anticathode and a diamond crystal. The ionization shows two maxima at about  $36.5^\circ$  and  $36.9^\circ$  which are due to two lines in the X-ray spectrum of rhodium. The small amount of ionization obtained at other angles is due to the continuous part of the X-ray spectrum.

Instead of using an ionization chamber the reflected rays may be received on a photographic plate and the angles found by measuring the

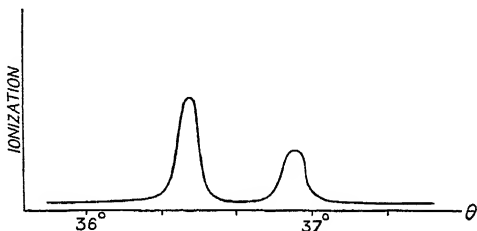


Fig. 13

distances between the images on the plate. The ionization method has the advantage that it gives the relative intensities of the lines accurately. Any particular line is only reflected when the crystal is set so that the glancing angle satisfies the equation

$$n\lambda = 2d \sin \theta.$$

To photograph all the lines which can be reflected from a crystal face it is therefore necessary to slowly rotate the crystal backwards and forwards during the exposure of the plate. In this way a photograph of all the lines reflected in the range of angles covered can be obtained. Each line can be reflected at all the angles given by  $n\lambda = 2d \sin \theta$ , with  $n = 1, 2, 3, 4, \&c.$   $n\lambda$  of course must be less than  $2d$ , since  $\sin \theta$  cannot be greater than 1. The lines are usually observed on both sides of the incident beam, and half the angle between the two positions is taken as the value of  $\theta$  so as to eliminate zero errors.

### 8. Example of Use of X-ray Spectrometer.

As a simple example of the use of the X-ray spectrometer to determine the structure of crystals and the wave-lengths of X-rays we will consider the reflection of the X-rays from a palladium anticathode by crystals of sodium and potassium chlorides. A full account of the subject is given in *X-rays and Crystal Structure*, by W. H. and W. L. Bragg.

Sodium chloride and potassium chloride can be obtained in cubical crystals the faces of which are specified by the indices (100), (010), (001). Besides these faces the faces (110) and (111) often appear. Fig. 14 shows these faces.

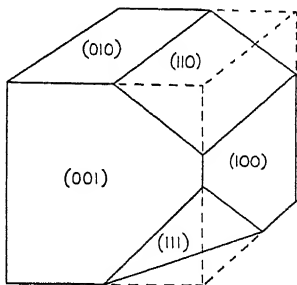


Fig. 14

The (100) face of KCl gives strong lines at  $5^\circ 23'$ ,  $10^\circ 49'$ ,  $16^\circ 20'$ . The sines of these angles are as 1 : 2 : 3, so that we conclude that they correspond to the same wave-length  $\lambda$  with  $n = 1, 2$ , and 3.

With the (110) and (111) faces the angles for  $n = 1$  are  $7^\circ 37'$  and  $9^\circ 23'$ . Now  $\sin 5^\circ 23' : \sin 7^\circ 37' : \sin 9^\circ 23' = 1 : \sqrt{2} : \sqrt{3}$  almost exactly. It appears therefore that the grating spaces for the faces (100), (110), and (111) are as  $1 : \frac{1}{\sqrt{2}} : \frac{1}{\sqrt{3}}$ . This is what we should obtain if the space lattice of the crystal consisted of equal cubes with an atom or group of atoms at each corner. The grating space for the (100) face would then be equal to the edge of a cube, for the (110) face to half the diagonal of a face of the cube and for the (111) face to one-third of the diagonal of the cube. These lengths are as  $1 : \frac{1}{\sqrt{2}} : \frac{1}{\sqrt{3}}$  in agreement with

the results obtained. We conclude that the crystals of KCl behave like a simple cubical space lattice.

The glancing angle for the same X-ray line from the (100) face of sodium chloride is  $6^\circ 0'$ . The grating space for this face is therefore smaller than for potassium chloride in the ratio  $\sin 5^\circ 23' : \sin 6^\circ 0' = 1 : 1.115$ . If the NaCl and KCl crystals have the same structure, which we should expect to be the case, then the grating spaces should be proportional to the cube roots of the molecular volumes.

The molecular volume of KCl is equal to the molecular weight  $39.10 + 35.46 = 74.56$  divided by its density 1.99 or 37.5 and that of NaCl is  $\frac{23.00 + 35.46}{2.163} = 27.02$  and  $\left(\frac{37.5}{27.0}\right)^{1/3} = 1.115$ , which agrees with the ratio of the grating spaces exactly.

The glancing angle for the (110) face of NaCl agrees with that for KCl in the same way, but for the (111) face of NaCl the glancing angle is just one-half that to be expected from the angle for KCl. In the case of NaCl the grating spaces for the faces (100), (110), and (111) are as  $1 : \frac{1}{\sqrt{2}} : \frac{2}{\sqrt{3}}$ , instead of  $1 : \frac{1}{\sqrt{2}} : \frac{1}{\sqrt{3}}$  as with

KCl. The numbers of electrons in K, Cl, and Na atoms are 19, 17, and 11 respectively, so that K and Cl atoms should be nearly equally efficient as scatterers of X-rays but Na atoms little more than half as efficient. Thus in a KCl crystal all the atoms are nearly equal as regards X-ray scattering, so that it makes little difference whether a K atom or a Cl atom is at any particular point. This suggests that in the simple cubical lattice of a KCl crystal the group is either one K atom or one Cl atom, and that in a NaCl crystal the arrangement of the atom is the same as in a KCl crystal, but that the effect on X-rays is not the same because of the difference between the number of electrons in Na and Cl atoms.

Since the grating spaces for the (100) and (110) faces correspond we conclude that the planes parallel to these faces contain equal numbers of metal and chlorine atoms in each case, whereas in the case of the (111) faces the planes contain all metal atoms and all chlorine atoms alternately. In the case of KCl the (111) grating space is the distance from a plane containing only K atoms to the next plane containing only Cl atoms, since these planes are practically equal as scatterers, but in the case of NaCl crystals the (111) grating space is the distance from a plane containing only Na atoms to the next plane containing only Na atoms and so is twice that corresponding to the (111) face in KCl. The arrangement of the atoms is therefore as shown in fig. 15. The black circles represent one sort of atom and the white circles the other sort. The atoms with equal numbers are all in the same (111) plane. Each sort of atom is arranged in a face-centred cubical lattice, that is, at the corners of cubes and in the middle of each face.

relative intensities of the reflected rays for different values of  $n$  in the equation  $n\lambda = 2d \sin\theta$ , or the relative intensities of the spectra of different orders, give information about the arrangement of the atoms. For example, it is that with the (111) face of sodium chloride the lines for which  $n = 2$  are stronger than those for which  $n = 1$ . The grating space in this case is the distance from a layer of Cl atoms to the next layer of Cl atoms. The rays reflected from the layers of Na atoms in between, when  $n = 1$ , are half a wave-length out of phase with the waves from the next layer of Cl atoms and so diminish the intensity of the reflected beam but, when  $n = 2$ , the rays from the Na atoms are in phase with the rays from the Cl atoms and so a strong reflected beam is obtained. In the case of potassium chloride the rays from the layers of K atoms are practically in phase with the rays from the layers of Cl atoms, so that if  $d$  is regarded as the distance from a Cl layer to the next Cl layer, then when  $n = 1$  there is practically a strong reflected beam, so that  $d$  is practically the distance from a Cl layer to the next Cl layer, as we have seen. These considerations serve to support the assumed arrangement of the atoms in these crystals.

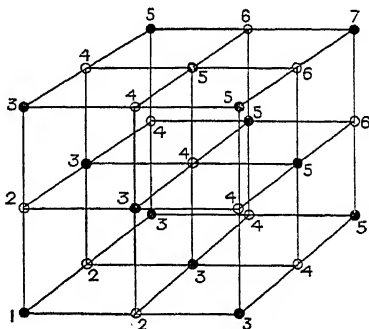


Fig. 15

When the arrangement of the atoms in a crystal has been decided, the absolute values of the grating spaces can be calculated from the known masses of the elements and the density of the crystal. The mass of a NaCl molecule is  $< 1.663 \times 10^{-24}$  grams, since  $1.663 \times 10^{-24}$  is the mass of one atom of sodium of atomic weight 1.0077. This gives  $9.797 \times 10^{-23}$  grams. The volume occupied by one atom of either sodium or chlorine in the crystal is therefore  $\frac{10^{-23}}{1.63} = 22.65 \times 10^{-24}$  c. c. The grating space for the (100) face is the edge of the elementary cubes, of which there is one to each atom,

$$d_{(100)} = (22.65 \times 10^{-24})^{1/3} = 2.83 \times 10^{-8} \text{ cm.}$$

The wave-length of the X-ray spectral line for which  $\theta = 6^\circ 0'$  when  $n = 1$  is

$$\lambda = 2 \times 2.83 \times 10^{-8} \times \sin(6^\circ 0') = 0.592 \times 10^{-8} \text{ cm.}$$

When the grating space for a face of a crystal has been found, the crystal can be used to determine the wave-lengths of X-ray spectral lines. The crystals usually used are calcite ( $d = 3.028 \times 10^{-8}$  cm.) or rock salt ( $d = 2.814 \times 10^{-8}$  cm.).

### Wave-lengths of Characteristic X-rays of the Elements. Moseley's Work.

Characteristic X-rays of any element are obtained by using it or one of its compounds as the anti-cathode of an X-ray tube such as the Coolidge tube.

The elements can also be made to emit their characteristic X-rays by exposing them to X-rays of sufficiently short wave-lengths.

The wave-lengths of the lines in the X-ray spectra of many of the

elements were first measured by Moseley, who discovered the very important fact that the frequencies vary in a regular way with the atomic number.

The X-ray spectral lines of any element can be arranged in series or groups of lines. The series having the highest frequencies is called the  $K$  series, that with the next highest frequencies the  $L$  series, and so on. The  $K$  series usually contains four strong lines known as  $K_{\alpha_2}$ ,  $K_{\alpha_1}$ ,  $K_{\beta}$ , and  $K_{\gamma}$ . The following table gives the wave-lengths of these  $K$  series lines in Ångstrom units ( $= 10^{-8}$  cm.) for several elements, and the atomic numbers of these elements. The atomic number of an element, except in one or two cases, is the number giving the position of the element in a list of the elements arranged in the order of increasing atomic weights. Thus the atomic numbers of hydrogen, helium, and lithium are 1, 2, and 3 respectively.

Element.	Atomic Number.	Wave-lengths of K Series.			
		$\alpha_2$ .	$\alpha_1$ .	$\beta$ .	$\gamma$ .
Sodium ..	11	—	11.8836	11.591	—
Potassium ..	19	3.738	3.7339	3.4464	—
Iron ..	26	1.932	1.9324	1.7540	1.736
Bromine ..	35	1.040	1.035	0.929	0.914
Rhodium ..	45	0.6164	0.6121	0.5453	0.5342
Cæsium ..	55	0.402	0.398	0.352	—
Tungsten ..	74	0.2134	0.2086	0.1842	0.17901

Moseley found that if the square roots of the frequencies of the  $K_{\alpha}$  lines are plotted against the atomic numbers then a curve is obtained which is very nearly straight. Approximately, if  $\nu_{K_{\alpha}}$  is the frequency,

$$\nu_{K_{\alpha}} = \frac{3}{4}R(N - a)^2,$$

where  $N$  is the atomic number,  $a$  is a constant, and  $R$  is a constant which is equal to the Rydberg constant of the formulæ for optical spectral series. For example, the frequencies of the series in the spectrum of atomic hydrogen are given by  $\nu = R\left(\frac{1}{n^2} - \frac{1}{m^2}\right)$ , where  $n$  and  $m$  are integers. For the strongest line of the  $L$  series, the  $L_{\alpha}$  line, Moseley found

$$\nu_{L_{\alpha}} = \frac{5}{36}R(N - b)^2.$$

Now  $\frac{3}{4} = \frac{1}{1^2} - \frac{1}{2^2}$  and  $\frac{5}{36} = \frac{1}{2^2} - \frac{1}{3^2}$ .

# 10. Quantum Theory of Spectra. Energy Levels. *K*, *L*, *M*, and *N* Series. Quantum Numbers.

According to Bohr's quantum theory, the frequencies of the spectral lines of an atom consisting of a nucleus with a positive charge  $Ne$  and a single electron are given (Chap. V, section 13) by

$$\nu = RN^2 \left( \frac{1}{n^2} - \frac{1}{m^2} \right).$$

Moseley's formulæ for the  $K_\alpha$  and  $L_\alpha$  lines are clearly closely related to this last equation, and it was immediately seen that Moseley's results indicate that the atomic numbers of the elements are equal to the nuclear charges expressed in terms of the protonic charge  $e$ . This has been confirmed by all subsequent work, and in particular by researches on the scattering of  $\alpha$ -rays.

According to the quantum theory of spectra the frequencies of the lines are given by

$$h\nu = W_m - W_n,$$

where  $h$  is Planck's constant, and  $W_m$  and  $W_n$  denote the energies of the atom in two of its possible states. The radiation is emitted when the atom changes from a state with energy  $W_m$  to one with less energy  $W_n$ , the change being supposed to consist of an electron dropping from an outer possible orbit to an inner vacant orbit. In the case of an atom having only one electron outside the nucleus we have

$$\frac{W_n}{h} = A - \frac{RN^2}{n^2},$$

so that 
$$\frac{W_m - W_n}{h} = RN^2 \left( \frac{1}{n^2} - \frac{1}{m^2} \right) = \nu.$$

For atoms with more than one electron outside the nucleus no theoretical expression for the energies of the possible states or energy levels of the electrons has been worked out, but since every frequency emitted gives the difference between two energy levels it is possible to deduce the energy levels of an atom from the frequencies of the lines which it emits.

The principal lines in the X-ray spectra of any element can be regarded as due to transitions of electrons between energy levels which are called the *K*, *L*, *M*, and *N* levels. The *K* series is due to transitions from the *L*, *M*, and *N* levels to the *K* level, the *L* series to transitions from the *M* and *N* levels to the *L* level, and the *M* series to transitions from the *N* level to the *M* level. Each of these levels except the *K* level is really not a single level but a group of levels which

do not differ much from each other. The number  $n$  in the equation

$$\frac{W_n}{h} = A - \frac{RN^2}{n^2}$$

is called the *quantum number* fixing the state of the atom, and the state is referred to as an  $n$ -quantum state, or an  $n$ -quantum energy level, of the electron. This terminology is carried over to the more complicated atoms, and the  $K$ ,  $L$ ,  $M$ , and  $N$  states or energy levels are referred to as one, two, three, and four quantum states or energy levels respectively.

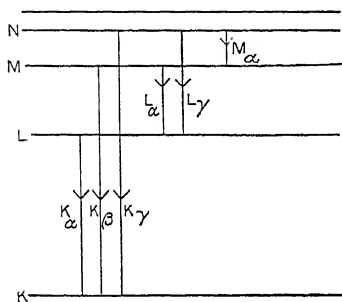


Fig. 16

The energy levels and the transitions between them are shown diagrammatically in fig. 16. The horizontal lines represent the energy levels, and the vertical distances between them the energy differences

such as  $W_L - W_K$ . The vertical distance between the  $K$  level and the line at the top is supposed to represent the energy required to remove an electron from the  $K$  level right out of the atom. The X-ray spectral lines are represented by the vertical lines, which show a transition from the level at the top of the line to that at the bottom. We see at once that

$$\nu_{K\alpha} + \nu_{L\alpha} = \nu_{K\beta}, \quad \nu_{K\alpha} + \nu_{L\gamma} = \nu_{K\gamma}, \quad \nu_{L\alpha} + \nu_{M\alpha} = \nu_{L\gamma},$$

with other similar relations. It is found that these relations between the frequencies are accurately true.

## 11. X-ray Absorption Spectra. Energy Levels.

The absorption spectrum of an element for X-rays can be observed with the X-ray spectrometer by passing the X-rays from a Coolidge tube through a plate of the substance, and examining the spectrum of the transmitted rays either photographically or with the ionization chamber. It is found that there are certain critical wave-lengths at which the absorption suddenly increases as the wave-length diminishes. A critical absorption wave-length characteristic of an element is a wave-length such that the element absorbs X-rays of shorter wave-length than the critical value more than it does wave-lengths longer than the critical value. If the element is present in a gas through which the X-rays are passed, the ionization of the gas is greater for X-rays with wave-lengths shorter than the critical value than for longer wave-lengths. The increased absorption is accompanied by increased ionization. According to the quantum theory, absorption

occurs when an electron in the atom is moved from its orbit or energy level to the free state outside the atom. In a normal atom all the energy levels are occupied by electrons, so that it is not possible for the radiation to move an electron from one level to another. If  $E_n$  denotes the energy required to move an electron from an  $n$ -quantum level to a position outside the atom, then the incident radiation can only move the electron if its frequency  $\nu$  is at least as great as that given by  $h\nu = E_n$ . When an electron drops back from just outside into the vacant level the atom emits a quantum of radiation  $h\nu = E_n$ . The critical absorption frequencies of an element therefore give the values of  $E_n$  for the element. The  $K$  series, for example, is due to electrons falling from the  $L$ ,  $M$ ,  $N$ , &c., levels into the  $K$  level. The frequency of the line in the  $K$  series of highest frequency should therefore be practically identical with the critical absorption frequency corresponding to the  $K$  level, for the highest energy level must nearly correspond to a point just outside the atom. It is found in fact that the  $K$  critical absorption frequencies are only a fraction of one per cent greater than the frequencies found for the  $K$  lines of highest frequency.

The critical absorption frequencies therefore give valuable information about the energy levels in the atom.

If  $W_0$  denotes the energy of a normal neutral atom which has all its energy levels occupied by electrons, and  $W_m$  the energy of the atom when one  $m$ -quantum level is vacant, then

$$W_m = W_0 + E_m.$$

If now an electron drops from an  $n$ -quantum level to the vacant  $m$ -quantum level the energy of the atom diminishes by

$$W_n - W_m = (W_0 + E_n) - (W_0 + E_m) = E_n - E_m = h\nu,$$

where  $\nu$  is the frequency of the radiation emitted. Since we are only concerned with energy differences, we may if we like take  $W_0$  to be zero and the energy of an atom with one  $m$ -quantum level vacant as equal to  $E_m$ , the work required to move an electron from an  $m$ -quantum level in the normal atom to just outside the atom. It is found that there is only one one-quantum or  $K$  level, but that there may be three  $L$  levels, five  $M$  levels, seven  $N$  levels, five  $O$  levels, and three  $P$  levels. These levels are found from the critical absorption frequencies when these have been observed, or from the frequencies of the lines in the different series. It is supposed, for example, that the three  $L$  levels correspond to electron orbits of different shapes but about the same size. The three  $L$  levels are numbered  $2(1, 1)$ ,  $2(2, 1)$ ,  $2(2, 2)$  respectively. The electrons in the normal atom are supposed therefore to be arranged in groups or sets corresponding to the groups of levels. This question is discussed in the chapter on Constitution of Atoms (Chapter XII).

## 12. Energy of Electrons and Frequency of Rays. De Broglie's Apparatus.

As we have seen, an element may be caused to emit its characteristic X-ray spectrum by bombarding it with electrons, as when it forms the anti-cathode of a Coolidge tube. The bombarding electrons do not excite a particular spectral line unless they have enough energy. For example, to cause the anti-cathode to emit its  $K$  series lines the bombarding electrons must have at least as much energy as is required to remove an electron from the  $K$  level outside the atom. When the  $K$  level is vacant, electrons may fall into it from any of the higher levels, so that all the  $K$  lines appear together as the energy of the bombarding electrons is gradually increased. If the potential difference between the cathode and anti-cathode is  $V$ , the energy of the bombarding

electrons is  $Ve$ , so that to excite a series of lines due to electrons falling into a vacant  $n$ -quantum level requires that  $Ve \geq E_n$ . It is found that these conclusions from the theory are in agreement with the facts.

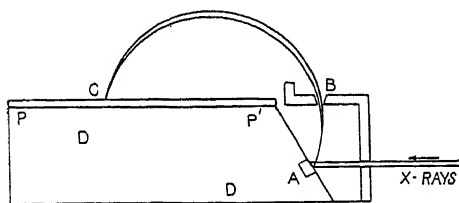


Fig. 17

The anti-cathode always emits a continuous X-ray spectrum as well as the spectral lines. It is found that this continuous spectrum ends sharply at a definite frequency, and no rays with a greater frequency than this limit are observed. The frequency at the limit is given by  $Ve = h\nu$ . This equation was shown to be accurately true by Duane and Hunt for values of  $V$  from 4500 to 170,000 volts.

When X-rays are passed through matter they cause the atoms to emit electrons. A quantum  $h\nu$  of the incident rays is absorbed by an electron. If  $E_n$  is the energy required to remove the electron just outside the atom from its energy level, the electron is shot out of the atom with kinetic energy equal to  $h\nu - E_n$ . By measuring the kinetic energies of the electrons shot out by rays of known frequencies the energy levels of the atoms can be determined. The results obtained in this way agree with those obtained by the other methods.

The apparatus used by De Broglie for measuring the energies of the electrons emitted by matter when acted on by X-rays is shown in fig. 17. DD is a heavy metal block into which a small piece of the substance to be investigated is put at A. A narrow beam of X-rays is allowed to fall on the substance at A as shown. Some of the electrons from A pass through a slit at B and are deflected by a uniform magnetic field perpendicular to the plane of the paper so that they describe

circular paths and fall on a photographic plate  $PP'$  at  $C$ . The whole apparatus is contained in a box which is exhausted, and the electrons are consequently not deflected by collisions with gas molecules. If  $m$  is the mass of an electron and  $v$  its velocity, then the radius  $R$  of its circular path is given by  $\frac{mv^2}{R} = Hev$ , where  $H$  is the strength of the magnetic field. We have also  $m = \frac{m_0}{\sqrt{1-v^2/c^2}}$ , where  $m_0$  is the mass when  $v = 0$ . By means of these equations  $v$  can be found, with the help of the known value of  $e/m_0$ . The kinetic energy of the rays is equal to

$$c^2 m_0 \left\{ \frac{1}{\sqrt{1-v^2/c^2}} - 1 \right\},$$

and so can be calculated.

It is found that a series of lines is obtained on the photographic plate, showing that the electrons come off with definite velocities in accordance with the quantum theory. The energies of the electrons are always equal to  $h\nu - E_n$ , where  $\nu$  is a frequency of the incident X-rays and  $E_n$  is the energy required to remove an electron from one of the energy levels in the atoms of the substance at  $A$ .

### 13. Scattering of X-rays. Classical Theory.

When a narrow beam of X-rays is passed through a thin plate of a substance of small atomic weight, in which the rays do not excite the characteristic X-rays of the substance, scattered X-rays are emitted by the plate in all directions. The intensity of these scattered rays in any direction can be measured by passing a narrow beam of them into an ionization chamber and measuring the conductivity they produce in it.

It is found that the intensity of the scattered rays varies with the angle between their direction and that of the incident beam. It is greatest when this angle is small, least at  $90^\circ$ , and greater near  $180^\circ$  than at  $90^\circ$ . If the intensity at  $90^\circ$  is taken as unity then that at  $180^\circ$  is about 2 and near  $0^\circ$  about 6.

If we assume the radiation to be scattered by free electrons according to the classical electromagnetic theory, it is easy to calculate the total energy of the scattered rays. On the classical theory (p. 15), an electron moving with an acceleration  $f$  radiates energy at the rate of  $\frac{1}{6\pi} \frac{e^2 f^2}{c^3}$  ergs per second. If  $F$  denotes the electric field strength in the incident X-rays then  $mf = Fe$  where  $m$  is the mass and  $e$  the charge of the electron. The rate of radiation is therefore  $\frac{e^4 F^2}{6\pi c^3 m^2}$ . The energy density in the incident rays is  $\frac{1}{2}(F^2 + H^2)$ , where  $H$  is the strength of the magnetic field. But  $F = H$ , and the energy density is therefore  $F^2$ .

Hence the energy flowing through unit area in unit time is  $c\bar{F}^2$ , where  $\bar{F}^2$  denotes the average value of  $F^2$  in the incident rays, and the energy scattered by the electron in unit time is

$$\frac{c^4 \bar{F}^2}{6\pi e^3 m^2}.$$

If a piece of matter containing  $N$  electrons is placed in the incident beam, the total energy of the rays scattered by it will be equal to  $Nc^4 \bar{F}^2 / 6\pi e^3 m^2$ . The ratio of the scattered energy to the energy flowing through unit area in the incident beam is therefore  $\frac{Nc^4}{6\pi m^2 c^4}$ . This ratio can be determined approximately by measuring the ionizations produced by the incident and scattered rays, and thus  $N$  can be found. In this way Barkla showed that the number of electrons in atoms of small atomic weight like carbon is not far from the atomic number. The classical theory thus appears to give results for light elements which agree at any rate roughly with the observed scattering.

#### 14. A. H. Compton's Quantum Theory of Scattering.

It is found that the wave-length of the scattered rays is not equal to that of the incident rays, as it should be according to the classical theory. This change of wave length by scattering has been investigated by A. H. Compton, who has shown that it can be explained on the quantum theory in a simple way.

We suppose that the electron receives a quantum of energy  $h\nu$  from the incident rays and emits a quantum  $h\nu'$  of scattered rays in a direction making an angle  $\theta$  with the direction of the incident rays. The momentum of the incident quantum is  $h\nu/c$  and that of the scattered quantum is  $h\nu'/c$ , so that the electron receives an impulse and starts off with a velocity  $v$  in a direction making an angle  $\phi$  with the direction of the incident rays. Compton supposes that energy and momentum are conserved, so that

$$\begin{aligned} h\nu &= h\nu' + mc^2 \left( \sqrt{1 + v^2/c^2} - 1 \right), \\ \frac{h\nu}{c} &= \frac{h\nu'}{c} \cos \theta + \cos \phi \cdot \sqrt{1 + v^2/c^2} \cdot m, \\ 0 &= -\frac{h\nu'}{c} \sin \theta + \sin \phi \cdot \sqrt{1 + v^2/c^2} \cdot m, \end{aligned}$$

where  $m$  is the mass of an electron at rest. Thus we have three equations to determine  $\nu'$ ,  $v$ , and  $\phi$ . Eliminating  $\phi$  from the last two equations we get

$$v^2 - 2\nu\nu' \cos \theta + \nu'^2 = \frac{m^2 c^2 v^2}{h^2 (1 + v^2/c^2)},$$

which with the first equation gives

$$\nu' = \nu / \{1 + \alpha(1 - \cos \theta)\}$$

where  $\alpha = h\nu/mc^2$ .

If  $\lambda$  is the wave-length of the incident rays and  $\lambda'$  that of the scattered rays, so that  $\nu\lambda = \nu'\lambda' = c$ , then

$$\frac{\lambda'}{\lambda} = 1 + \alpha(1 - \cos \theta)$$

$$\text{or} \quad \lambda' - \lambda = (h/mc)(1 - \cos \theta).$$

Thus this theory indicates that  $\lambda'$  is greater than  $\lambda$  unless  $\theta = 0$ . The difference  $\lambda' - \lambda$  is independent of  $\lambda$ , and the percentage change is therefore large for small wave-lengths but small for large ones. Putting in the known values of  $h$ ,  $m$ , and  $c$  we get

$$\lambda' - \lambda = 0.0242(1 - \cos \theta),$$

where  $\lambda' - \lambda$  is in Ångstrom units ( $1 = 10^{-8}$  cm.). Solving the equations for the angle  $\phi$ , we get

$$\tan \phi = \frac{\cot(\theta/2)}{1 + \alpha},$$

and for the kinetic energy of the electron we get

$$2mc^2\alpha^2 \cos^2 \phi / (1 + 2\alpha + \alpha^2 \sin^2 \phi).$$

The electrons which receive energy from the X-rays in this way are called recoil electrons to distinguish them from photo-electrons, which receive energy  $h\nu$ .

A. H. Compton in 1923 scattered X-rays from a molybdenum anti-cathode by a carbon plate and measured the wave-lengths of the scattered rays with a Bragg spectrometer. It was found that the lines in the spectrum of the scattered rays were usually doublets, each doublet consisting of an unmodified line of wave-length  $\lambda$  and a modified line of wave-length  $\lambda'$ . The difference  $\lambda' - \lambda$  was approximately equal to  $0.0242(1 - \cos \theta)$  in agreement with the quantum theory. This result has since been confirmed by several other observers.

## 15. Experiments supporting Quantum Theory of X-rays.

The quantum theory of the scattering of X-rays seems to require that the X-rays consist of particles having energy  $h\nu$  and momentum  $h\nu/c$ . We may suppose that these particles execute some kind of oscillation of frequency  $\nu$  as they move along with the velocity of light  $c$ . It is difficult to see how phenomena such as diffraction of X-rays by crystals, which apparently depend on interference between trains of waves, can be explained on this theory.

Some very interesting experiments have been made by Bothe and Geiger, and by Compton and Simon, which strongly support the particle or quantum theory of X-rays. Bothe and Geiger passed a narrow beam of X-rays through hydrogen. Two ionization chambers were arranged on either side of the beam in such a way that if a scattered quantum entered one chamber the scattering electron according to the theory would enter the other. The ionization in the chambers was greatly increased by ionization by collisions so that the effect due to a single quantum or electron could be detected. According to the theory we should expect that when ionization was produced in one chamber ionization would be produced at the same instant in the other. It was found that simultaneous ionizations occurred in about 10 per cent of the observed cases, which Bothe and Geiger considered to be considerably more than could be explained by chance coincidences.

Compton and Simon passed a narrow beam of X-rays of very short wave-length into a C. T. R. Wilson cloud chamber, and examined the electron tracks produced. The recoil electrons produce short tracks and the photo-electrons or  $\beta$ -rays emitted by atoms with energy approximately  $h\nu$  produce much longer tracks, so that it is easy to distinguish between the two. It was found that occasionally a recoil track started from the path of the X-ray beam and simultaneously a  $\beta$ -ray track from a neighbouring point outside the beam. The  $\beta$ -ray track was assumed to be produced by an electron shot out of an atom by the scattered quantum which produced the recoil track. The angle  $\phi$  between the initial direction of the recoil track and the X-rays was observed, and also the angle  $\theta$  between the X-ray beam and the direction from the beginning of the recoil track to the beginning of the  $\beta$ -ray track. It was found that the values of  $\phi$  observed agreed approximately with those calculated by means of the equation

$$\tan \phi = \frac{\cot(\theta/2)}{1 + \alpha}.$$

These results and those of Bothe and Geiger seem to support very strongly the particle theory of X-rays and to be quite inexplicable on the wave theory. It appears that there is a large number of facts which agree with the classical wave theory of light and X-rays and also a large number of facts which agree with a particle or quantum theory. Each set of facts agrees with one theory and seems definitely inconsistent with the other theory. Since agreement between a theory and a set of facts does not prove the theory true, whereas disagreement does prove the theory untrue or at least inadequate, the proper conclusion to be drawn is clearly that both theories are inadequate or untrue. Presumably a new theory will eventually be discovered which will be capable of explaining all the facts.

16.  $\gamma$ -rays.

The most penetrating rays emitted by radioactive bodies are called  $\gamma$ -rays and are found to have properties similar to those of very penetrating X-rays. They are not deflected by electric or magnetic fields and carry no charge of electricity. They affect a photographic plate and ionize gases like X-rays. The  $\gamma$ -rays from 30 mgm. of radium can be detected, after passing through 30 cm. of iron, by the ionization they produce.  $\gamma$ -rays are only emitted by radioactive bodies which also emit  $\beta$ -rays. This suggests that  $\gamma$ -rays are produced by  $\beta$ -rays or vice versa. It has been shown that  $\beta$ -rays produce  $\gamma$ -rays when they are stopped by lead, just as X-rays are produced by the impact of high-velocity electrons on the anti-cathode of an X-ray tube. It is also found that  $\gamma$ -rays produce  $\beta$ -rays when they are absorbed by matter, just as X-rays cause the emission of  $\beta$ -rays. Rutherford showed that when  $\beta$ -rays are produced by  $\gamma$ -rays, which were produced by  $\beta$ -rays from a radioactive body, then the  $\beta$ -rays produced by the  $\gamma$ -rays have the same energy as the  $\beta$ -rays which produced the  $\gamma$ -rays.

The spectra of the  $\gamma$ -rays from radioactive bodies have been examined by means of the Bragg crystal spectrometer or similar apparatus, and it is found that the spectra contain lines having wave-lengths characteristic of the elements emitting the rays. The wave-lengths range from  $0.07 \times 10^{-8}$  cm. to  $0.4 \times 10^{-8}$  cm. The  $\gamma$ -rays from some radioactive bodies appear to belong to the *K* or *L* series of X-rays, and so must be due to an electron falling from an outer level in the atom into the *K* or *L* level. The  $\beta$ -rays emitted by the radioactive nucleus may be supposed to knock an electron out of the *K* or *L* level and the  $\gamma$ -rays to be emitted when an electron falls into the place left vacant.

When  $\gamma$ -rays are scattered by light elements like carbon, modified rays having a longer wave-length are obtained as with X-rays. The change of wave-length appears to agree with that calculated on Compton's theory of X-ray scattering.

## 17. The J Phenomenon.

The scattering of X-rays has been studied very extensively by Barkla, who finds results which apparently are not in agreement with Compton's quantum theory. The following summary of Barkla's results is quoted from one of his recent papers. (1) When a heterogeneous X-radiation is scattered, the scattered radiation has either precisely the same absorbability as the primary, or there is a well-marked difference between the absorbabilities when measured in any one substance. (2) When scattered radiation is different from the primary when measured in certain substances, its absorbability may be, and frequently is, precisely like that of the primary when measured

in certain other substances. (3) Even after transmission through substances which show differences between primary and secondary radiations, there is still no difference between the two when measured in certain other substances. (4) When the scattered radiation is observed in different directions making angles of  $30^\circ$ ,  $60^\circ$ , and  $90^\circ$  with the primary beam, and when these radiations differ from the primary in their absorbability in a certain substance, they in general differ by precisely the same amount. When a difference appears with increasing angle of scattering it takes place abruptly by a jump.

The changes in absorbability are referred to by Barkla as the J phenomenon. He considers that a heterogeneous beam of X-rays has properties depending on the distribution of energy in its spectrum which cannot be deduced from the properties of the beams of homogeneous rays of which the heterogeneous beam may be regarded as composed. He considers that his results do not agree with either the classical wave theory or the quantum particle theory of X-rays.

Barkla's results seem to show that absorbability does not define the properties of a heterogeneous beam of X-rays. The absorbabilities of two beams may be the same in some different substances and different in other substances.

#### 18. Millikan's Cosmic Rays.

It has been known for a long time that there is present at the earth's surface a small amount of highly penetrating radiation which produces ionization in gases and is probably similar in nature to  $\gamma$ -rays. A gold-leaf electroscope slowly loses its charge, even when the leak along the support of the gold leaf is prevented, showing that the air in the electroscope is being ionized to a small extent. It is found that surrounding the electroscope by a thick layer of lead diminishes the ionization in it, showing that some of the ionization is due to penetrating radiation. McLennan found that the ionization in an electroscope could be diminished by sinking it to a considerable depth in Lake Ontario.

Even when all the penetrating radiation coming in from outside is cut off in this way some ionization remains, which may be attributed to the radiation from minute traces of radioactive bodies in the walls of the electroscope. The ionization may be reduced to only 2 or 3 ions per cubic centimetre per second in this way. It is found that the ionization in an electroscope increases when the electroscope is taken up in a balloon, which suggests that there may be some penetrating radiation coming downwards from the upper regions.

This question has recently been carefully investigated by Millikan. Millikan measured the ionization in an electroscope at different levels on mountains in California, and also the ionization in the electroscope when sunk to different depths in lakes at different levels in the moun-

tains. He found that the ionization due to penetrating radiation produces 1.4 pairs of ions per cubic centimetre per second at sea-level, 2.6 at 1600 m., 4.8 at 3600 m., and 5.9 at 4300 m. above sea-level.

On sinking the electroscope in Muir Lake, which is 11,800 ft. above sea-level, the ionization dropped from 13.3 ions per cubic centimetre per second at the surface of the lake to 3.6 at 50 ft. below the surface, and remained constant at greater depths. In Arrowhead Lake at 5100 ft. it dropped from 7 at the surface to a minimum of 3.6 at depths greater than about 40 ft. The 6700 ft. of air between the levels of the two lakes is equal in weight to about 6 ft. of water, and it was found that the ionizations in Muir Lake were equal to those in Arrowhead Lake at depths 6 ft. greater in Muir Lake than in Arrowhead Lake.

In both lakes the ionization diminished with depth, so that the difference between the ionization at a depth  $x$  and that at great depths (3.6) was roughly proportional to  $e^{-\mu x}$  where  $\mu$  is a constant. The values of  $\mu$  found are from 0.18 per metre of water to 0.3 per metre of water. It was found that the intensity of the penetrating radiation was the same at all times, night and day, and it was concluded that the rays come into the atmosphere from the surrounding space equally in all directions. The wave-length of these cosmic rays as estimated from their absorption co-efficient is about  $0.0005 \times 10^{-8}$  cm., which is much shorter than that of any previously known rays. Millikan obtained evidence of the existence of high-velocity  $\beta$ -rays produced by the cosmic rays. It seems possible that these cosmic rays may be produced by radioactive changes in the small amount of matter supposed to be diffused throughout interstellar space. However, it is not clear why this matter should emit such rays if the matter on the earth does not do so.

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## CHAPTER VIII

# Optical Spectra

### 1. Principal, Sharp, Diffuse, and Fundamental Series.

The spectra of atomic hydrogen and of ionized helium are discussed in the chapter on quantum theory, and X-ray spectra in the chapter on X-rays. In the present chapter the series of spectral lines found in the optical spectra of many elements will be considered.

About 1890 Kayser and Runge discovered series of lines in the spectra of the alkali metals, and soon after this Rydberg showed that the frequencies \* of these lines could be expressed approximately as the differences of two *terms*, each term being of the form

$$\frac{N}{(n + \alpha)^2},$$

where  $N$  is a constant having the same value in all the series,  $\alpha$  a small constant, and  $n$  an integer. Rydberg's formula is analogous to Balmer's formula for the frequencies in the spectrum of atomic hydrogen, which are equal to the differences between two terms each of which is  $N/n^2$ .

Ritz showed that a more exact expression for the frequencies is got by taking the difference between two terms of the form  $\frac{N}{(n + \alpha + \beta/n)^2}$  or  $\frac{N}{(n + \alpha + \beta/n^2)^2}$ , where  $\alpha$  and  $\beta$  are small constants.

A particular *series* is represented by

$$\bar{\nu} = N \left\{ \frac{1}{(n' + \alpha)^2} - \frac{1}{(n + \alpha)^2} \right\},$$

where  $n'$  has the same integral value in all the lines of the series and  $n = n' + 1, n' + 2, n' + 3, \&c.$  When  $n$  is very large  $(n + \alpha)^{-2} = 0$ , and we get as the limit of the series  $N/(n' + \alpha)^2$ . The lines in a series

\* In practical spectroscopy, the word *frequency* is habitually used in the sense of "number of wave-lengths per centimetre". The word *wave-number* is sometimes used with the same meaning. To prevent the possibility of confusion, frequency in this conventional sense will be denoted here by the symbol  $\bar{\nu}$ , and frequency in the strict sense by  $\nu$  as hitherto. The constant  $N$ , called the Rydberg constant, has the value  $109732\text{cm}^{-1}$ . The constant  $K$  of Chap. V, section 13, is nearly equal to  $Nc$  and  $\nu = c\bar{\nu}$ .

get fainter as the limit is approached. Four such series have been found in the arc spectra of the alkali metals and alkaline earth metals. These series are known as the principal series, the sharp series, the diffuse series, and the fundamental series. The formulæ usually used for the frequencies in these four series are:

$$\text{Principal: } P(n) = P(\infty) - N/(n + P)^2, \quad n = 2, 3, 4 \dots,$$

$$\text{Sharp: } S(n) = S(\infty) - N/(n + S)^2, \quad n = 2, 3, 4 \dots,$$

$$\text{Diffuse: } D(n) = D(\infty) - N/(n + D)^2, \quad n = 3, 4, 5 \dots,$$

$$\text{Fundamental: } F(n) = F(\infty) - N/(n + F)^2, \quad n = 4, 5, 6 \dots,$$

where e.g.  $P(n)$  is the frequency of a line in the  $P$  series, and  $P$ ,  $S$ ,  $D$ ,  $F$  are the values of the constant  $\alpha$  in the respective series. In many cases each line of a series is not a single line but a doublet or triplet. Each series according to the above formulæ is determined by two constants, such as  $P(\infty)$  and  $P$ , which are different in different series. It was shown by Rydberg that these constants for different series are related in a simple way. He found that

$$S(\infty) = D(\infty) = \frac{N}{(2 + P)^2},$$

$$\text{and also that} \quad P(\infty) = \frac{N}{(1 + S)^2}.$$

From this it follows that  $P(\infty) - S(\infty)$  is equal to

$$P(\infty) - \frac{N}{(2 + P)^2} = P(2).$$

Runge pointed out that  $D(\infty) - F(\infty) = D(3)$ ,

$$\text{or} \quad F(\infty) = \frac{N}{(3 + D)^2}.$$

The four series may therefore be represented by

$$P(n) = \frac{N}{(1 + S)^2} - \frac{N}{(n + P)^2}, \quad n = 2, 3, 4 \dots,$$

$$S(n) = \frac{N}{(2 + P)^2} - \frac{N}{(n + S)^2}, \quad n = 2, 3, 4 \dots,$$

$$D(n) = \frac{N}{(2 + P)^2} - \frac{N}{(n + D)^2}, \quad n = 3, 4, 5 \dots,$$

$$F(n) = \frac{N}{(3 + D)^2} - \frac{N}{(n + F)^2}, \quad n = 4, 5, 6 \dots,$$

so that only four constants  $S$ ,  $P$ ,  $D$ , and  $F$  are required besides  $N$ .

These formulæ are usually written in an abbreviated form as follows:

$$\begin{aligned}P(n) &= 1S - nP, \\S(n) &= 2P - nS, \\D(n) &= 2P - nD, \\F(n) &= 3D - nF.\end{aligned}$$

Thus  $D(3)$  means the spectral line in the  $D$  series for which  $n=3$ , but  $3D$  stands for the term  $\frac{N}{(3+D)^2}$ .

## 2. The Combination Principle.

Rydberg and Ritz pointed out that spectral lines with frequencies given by the differences between any possible pair of the terms  $nP$ ,  $nS$ ,  $nD$ , and  $nF$  might be expected to appear. For example,  $2S - nP$ ,  $3S - nP$ , and so on. A great many such lines have been discovered, so that according to this principle, known as the combination principle, lines may occur with frequencies given by the abbreviated expression  $nX - n'X'$ , where  $n$  and  $n'$  are integers, and  $X$  and  $X'$  stand for  $P$ ,  $S$ ,  $D$ , or  $F$ . This result may be compared with the formula for the frequencies of the lines in the spectrum of atomic hydrogen,

$$\bar{\nu} = \frac{N}{n^2} - \frac{N}{n'^2}, \text{ where } n \text{ and } n' \text{ are integers.}$$

It appears as though spectral lines could be arranged in an infinite number of series each having an infinite number of terms. Of course only a comparatively small number of the lines indicated by these formulæ have been actually observed.

According to the quantum theory the terms like  $nP$ ,  $nS$ , &c., represent energy levels in the atoms. When an electron falls from one energy level to another having less energy, radiation is emitted of frequency  $\nu$  given by  $h\nu = E_1 - E_2$ . The line  $P(3)$  given

by  $P(3) = \frac{N}{(1+S)^2} - \frac{N}{(3+P)^2}$ , for example, is supposed to be emitted when an atom changes from a stationary state having energy

$C - \frac{Nhc}{(3+P)^2}$ , where  $C$  is a constant, to another state having energy

$C - \frac{Nhc}{(1+S)^2}$ , so that the frequency of the radiation emitted is given by

$$\bar{\nu} = P(3) = \frac{E_1 - E_2}{hc} = \frac{C}{hc} - \frac{N}{(3+P)^2} - \left( \frac{C}{hc} - \frac{N}{(1+S)^2} \right),$$

or

$$\bar{\nu} = P(3) = \frac{N}{(1+S)^2} - \frac{N}{(3+P)^2}.$$

### 3. Spectra of Atoms which have lost Electrons.

By using sparks between electrodes very near together in a very perfect vacuum and working with very large potential differences, Millikan and Bowen have obtained the spectra of many atoms which have lost one or more electrons, thus extending work of this kind initiated by Fowler, Paschen, and others.

We should expect atoms containing the same number of electrons but different nuclear charges to give similar spectra. Increasing the nuclear charge must diminish the distance of the electrons from the nucleus and so increase the energy of the atom, but it should not alter the arrangement of the electrons or the relative sizes of their orbits. Thus increasing the nuclear charge, keeping the number of electrons in the atom constant, should shift the whole spectrum towards shorter wave-lengths without altering the arrangement of the lines in it. In the case of atoms having only one electron we should expect the frequencies to be given by

$$\bar{\nu} = Z^2 N \left( \frac{1}{n^2} - \frac{1}{n'^2} \right),$$

where  $Z$  is equal to the number of ionic charges in the nucleus, or the atomic number, as in the cases of atomic hydrogen and ionized helium.

### 4. Spectra of Atoms containing 1, 2, 3, and 4 Electrons.

If we indicate the number of electrons lost by an atom by a suffix, so that, for example,  $O_{III}$  indicates an oxygen atom which has lost three electrons, then the atoms

$$H, He_I, Li_{II}, Be_{III}, B_{IV}, C_V, N_{VI}, O_{VII}, F_{VIII},$$

which all contain only *one* electron, should all give spectra represented by the above equation with  $Z = 1$  for  $H$ ,  $Z = 2$  for  $He_I$ ,  $Z = 3$  for  $Li_{II}$ , and so on. Lines given by  $\bar{\nu} = Z^2 N \left( \frac{1}{n^2} - \frac{1}{n'^2} \right)$  have so far only been observed in the spectra of  $H$ ,  $He$ , and  $Li$ .

The following atoms all contain *two* electrons:  $He$ ,  $Li_I$ ,  $Be_{II}$ ,  $B_{III}$ ,  $C_{IV}$ ,  $N_V$ ,  $O_{VI}$ ,  $F_{VII}$ . The spectra of these atoms should contain series represented approximately by

$$\bar{\nu} = Z^2 N \left\{ \frac{1}{(n' + a')^2} - \frac{1}{(n + a)^2} \right\},$$

where  $Z$  is equal to the atomic number, since such series have been found in the spectra of helium and  $Be_{II}$ .

The following atoms all contain *three* electrons:  $Li$ ,  $Be_I$ ,  $B_{II}$ ,  $C_{III}$ ,  $N_{IV}$ ,  $O_V$ ,  $F_{VI}$ .

The spectrum of Li contains series represented by

$$\bar{\nu} = \frac{N}{(n' + \alpha')^2} - \frac{N}{(n + \alpha)^2},$$

so we should expect similar series, with  $Z^2N$  instead of  $N$ , in the spectra of the other atoms with three electrons. Such series have been found by Fowler, Paschen, Millikan and Bowen, and others.

The following table given by Millikan and Bowen gives the values of the series terms for Li, Be<sub>I</sub>, B<sub>II</sub>, and C<sub>III</sub>. The difference between two such terms for the same element gives the frequency of a spectral line.

$n =$ $109732/n^2 =$	2. 27,433	3. 12,192.8	4. 6,858.4	5. 4,389.4
$P_1$ terms. $\begin{cases} \text{Li} \\ \text{Be}_I \div 4 \\ \text{B}_{II} \div 9 \\ \text{C}_{III} \div 16 \end{cases}$	$\begin{cases} 28,582.5 \\ 28,736.3 \\ 28,616.1 \\ 28,465.3 \end{cases}$	$\begin{cases} 12,560.4 \\ 12,596.2 \\ — \\ 12,504.3 \end{cases}$	$\begin{cases} 7,018.2 \\ 7,030.1 \\ — \\ — \end{cases}$	$\begin{cases} 4,473.6 \\ 4,477.6 \\ — \\ — \end{cases}$
$D$ terms. $\begin{cases} \text{Li} \\ \text{Be}_I \div 4 \\ \text{B}_{II} \div 9 \\ \text{C}_{III} \div 16 \end{cases}$	$\begin{cases} — \\ — \\ — \\ — \end{cases}$	$\begin{cases} 12,203.1 \\ 12,206.9 \\ 12,207.8 \\ 12,208.3 \end{cases}$	$\begin{cases} 6,863.5 \\ 6,865.1 \\ — \\ — \end{cases}$	$\begin{cases} 4,389.6 \\ 4,393.7 \\ — \\ — \end{cases}$
$F$ terms. $\begin{cases} \text{Li} \\ \text{Be}_I \div 4 \\ \text{B}_{II} \div 9 \end{cases}$	$\begin{cases} — \\ — \\ — \end{cases}$	$\begin{cases} — \\ — \\ — \end{cases}$	$\begin{cases} 6,856.1 \\ 6,858.8 \\ 6,860.2 \end{cases}$	$\begin{cases} 4,381.8 \\ 4,389.5 \\ — \end{cases}$

The second row of this table gives the values of  $N = 109,732$  divided by  $n^2 = 4, 9, 16$ , or  $25$ . These are the values of the terms in the equation  $\bar{\nu} = \frac{N}{n'^2} - \frac{N}{n^2}$ , which gives the frequencies of the spectral lines of atomic hydrogen.

The remaining rows give the values of the series terms deduced from the observed frequencies of the lines of Li, Be, B, and C atoms. The terms attributed to Be<sub>I</sub> are divided by 4, those attributed to B<sub>II</sub> by 9, and those attributed to C<sub>III</sub> by 16.

We see that the resulting quotients are nearly equal to the values of  $N/n^2$  and are nearly the same for the different atoms. Thus, for example, the spectrum of B<sub>II</sub> contains lines having nearly nine times the frequencies of lines in the spectrum of Li.

It is supposed that in these atoms, which contain three electrons, two electrons describe small orbits near the nucleus and the third a much larger orbit. The outer electron has a number of possible orbits

according to the Bohr theory, and lines with frequencies nearly equal to  $Z^2N\left(\frac{1}{n'^2} - \frac{1}{n^2}\right)$  are emitted when the outer electron drops from one outer orbit to another farther in. The nucleus and the two inner electrons form a small group having a total charge  $(Z - 2)e$  where  $Z$  is the atomic number, so that so long as the orbits of the outer electron do not come near this group the orbits are very similar to those of a single electron moving near a nucleus with charge  $(Z - 2)e$ .

For        Li,  $Z = 3$ , so that  $(Z - 2)e = e$ ;  
 for        Be,  $Z = 4$ , so that  $(Z - 2)e = 2e$ ;  
 for        B,  $Z = 5$ , so that  $(Z - 2)e = 3e$ ;  
 and for    C,  $Z = 6$ , so that  $(Z - 2)e = 4e$ .

Thus the appropriate constants for Li, Be<sub>I</sub>, B<sub>II</sub>, and C<sub>III</sub> are  $N$ ,  $4N$ ,  $9N$ , and  $16N$ .

The following atoms all contain *four* electrons:

Be, B<sub>I</sub>, C<sub>II</sub>, N<sub>III</sub>, O<sub>IV</sub>, F<sub>V</sub>.

These atoms have two outer electrons and should therefore give series represented by

$$\frac{Z^2N}{(n' + \alpha')^2} - \frac{Z^2N}{(n + \alpha)^2},$$

since such series are found in the spectra of Ca, Sr, Ba, and Ra, which all have two outer electrons.

### 5. Spectra of Atoms with 11 Electrons.

The following atoms all contain 11 electrons:

Na, Mg<sub>I</sub>, Al<sub>II</sub>, Si<sub>III</sub>, P<sub>IV</sub>, S<sub>V</sub>.

Sodium is supposed to have 2 electrons very near the nucleus, 8 farther out, and 1 outer electron which describes orbits usually well outside the inner group consisting of the nucleus and 10 electrons. The electrons in the other atoms with 11 electrons should be arranged in the same way, so that we should expect all these atoms to give series containing lines represented by

$$Z^2N\left(\frac{1}{(n' + \alpha')^2} - \frac{1}{(n + \alpha)^2}\right).$$

This is found to be the case. The following table gives the term values for these atoms divided in each case by  $(Z - 10)^2$ ,  $Z$  being the atomic number.

$n =$ $N/n^2 =$		3. 12,192.78	4. 6,858.44	5. 4,389.40	6. 3,048.19
S	Na/1	41,499.0	15,709.5	8,248.3	5,077.3
	Mg/4	30,316.9	12,865.6	7,120.3	4,517.3
	Al/9	25,494.89	11,476.82	6,535.29	—
	Si/16	22,756.83	10,633.65	6,168.72	—
	P/25	20,979.65	10,061.63	5,914.35	—
	S/36	19,729.56	9646.22	—	—
P	Na/1	24,475.7	11,176.1	6,406.3	4,151.3
	Mg/4	21,376.6	10,154.0	5,949.6	3,909.2
	Al/9	19,504.01	9,526.85	5,664.93	—
	Si/16	18,272.62	9,105.53	5,471.16	—
	P/25	17,401.88	8,802.20	—	—
	S/36	16,753.64	—	—	—
D	Na/1	12,276.2	6,900.4	4,412.5	3,061.9
	Mg/4	12,444.3	6,988.8	4,461.6	3,091.6
	Al/9	12,611.0	7,074.3	4,508.72	3,119.96
	Si/16	12,733.65	7,131.84	4,539.22	—
	P/25	12,811.8	7,164.04	—	—
	S/36	12,856.59	—	—	—
	Na/1	—	6,860.4	4,390.4	3,043.0
	Mg/4	—	6,866.8	4,394.3	3,051.2
	Al/9	—	6,871.28	4,397.61	3,053.83
	Si/16	—	6,874.19	4,399.97	3,055.97
	P/25	—	6,876.38	4,401.46	—
	S/36	—	6,878.11	—	—
F''	Na/1	—	—	4,388.8	—
	Mg/4	—	—	—	3,048.7
	Al/9	—	—	4,391.80	3,050.30
	Si/16	—	—	4,392.31	3,050.84
	P/25	—	—	4,392.74	3,051.14
	S/36	—	—	4,393.32	—
F'''	Na/1	—	—	—	3,046.3
	Mg/4	—	—	—	—
	Al/9	—	—	—	3,049.64
	Si/16	—	—	—	3,049.81
	P/25	—	—	—	3,050.21

This table is taken from a paper by Bowen and Millikan (*Physical Review*, March, 1925). The values of the series terms in it are due to Fowler, Paschen, and Millikan and Bowen.

It will be seen that the quotients of the *D* and *F* terms are nearly equal to the corresponding values of  $N/n^2$ . This is interpreted, as before, to mean that the orbits of the outer electron involved are well outside the inner orbits. Similar results to the above have been obtained with several other sets of atoms. We may remark, for example,

that potassium can be made to give a spectrum very similar to that of argon. The normal K atom has 19 electrons and the argon atom 18. It is therefore supposed that the K spectrum which resembles that of argon is due to K atoms which have lost one electron.

#### 6. Analogy with Moseley's Law. Second Quantum Numbers.

These results on optical spectra are closely analogous to Moseley's law for X-rays. Moseley found that the frequencies of the  $K_\alpha$  lines in X-ray spectra are given by

$$\bar{\nu} = \left(1 - \frac{1}{2^2}\right) N (Z - a)^2,$$

so that  $\sqrt{\bar{\nu}}$  is proportional to  $Z - a$ , where  $Z$  is the atomic number and  $a$  is a constant. The square roots of the values of the series terms of which the quotients by  $(Z - 10)^2$  are given in the above table are nearly proportional to  $Z - 10$ , except in the cases of the terms near the top of the table. This result is analogous to Moseley's law with  $a = 10$ .

Millikan and Bowen have pointed out that Moseley's law applies to the optical spectra of many other similar series of atoms.

The quantum theory of the fine structure of the atomic hydrogen and ionized helium lines is discussed in Chapter V. It appears that two integers or quantum numbers  $n_1$  and  $n_2$  are required to define the electron orbits, and that the series terms are approximately equal to  $NZ^2/(n_1 + n_2)^2$ . The major axis  $a$  of the orbit is determined by  $n_1 + n_2$  and the minor axis  $b$  by the equation  $\frac{b}{a} = \frac{n_1}{n_1 + n_2}$ .  $n_1$  is of course equal to or less than  $n_1 + n_2$ , but the value  $n_1 = 0$  is not admissible because the electron cannot be supposed to pass through the nucleus.

It is supposed that similar principles apply to the orbits of the electrons in other atoms besides H and He. Each orbit or atomic energy level is therefore supposed to be characterized by a principal quantum number and a second number  $n_1$ , which determines the shape of the orbit. The energy levels are not the same for orbits with the same principal number but different second quantum numbers. It is supposed that the terms of the different series are determined by the values of the second quantum number  $n_1$  as well as by the principal quantum number. For an  $S$  term  $n_1 = 1$ , for a  $P$  term  $n_1 = 2$ , for a  $D$  term  $n_1 = 3$ , and for an  $F$  term  $n_1 = 4$ .

For example, the principal series given by  $P(n) = 1S - nP$  is due to transitions from  $P$  levels for which  $n_1 = 2$  to the  $1S$  level for which  $n_1 = 1$ . We should expect  $n$  in the terms  $nP$  to be equal to the sum of the two quantum numbers  $n_1 + n_2$ , since  $nP = \frac{Z^2 N}{(n + P)^2}$ , and for

atomic hydrogen the terms are nearly equal to  $\frac{N}{(n_1 + n_2)^2}$ , so that  $n + P$  corresponds to  $n_1 + n_2$ . In terms which are nearly equal to  $\frac{Z^2 N}{n^2}$  it is clear that  $n = n_1 + n_2$ , but in other cases the value of  $n_2$  may not be known.

According to Sommerfeld only such transitions between  $S$ ,  $P$ ,  $D$ , and  $F$  terms are ordinarily possible as involve a change of  $\pm 1$  in the second quantum number  $n_1$ . This selection principle was derived by Sommerfeld in the case of atoms having only one electron by applying the principle of the conservation of momentum to the atom and the radiation emitted. It is assumed to apply to more complicated atoms also. The values of  $n_1$  assigned to the  $S$ ,  $P$ ,  $D$ , and  $F$  terms are in conformity with this selection principle.

### 7. The Series Terms for Sodium.

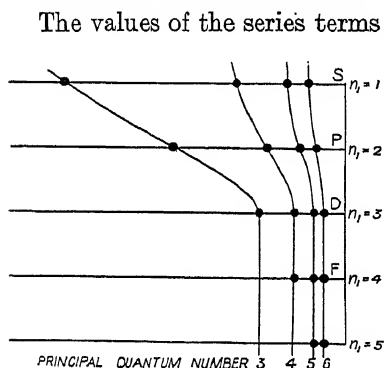


Fig. 1

The values of the series terms for sodium are shown graphically in the accompanying figure due to Bohr (fig. 1). The distances of the black dots from the vertical line on the right are proportional to the values of the series terms. The  $S$  terms are in the top row, the  $P$  terms in the second row, and so on. The curved lines are drawn through terms having the same principal quantum number, and the distances of these lines from the vertical line on the line for  $n_1 = 5$  are proportional to the

corresponding terms of a hydrogen atom.

We see that the  $D$ ,  $F$ , and  $n_1 = 5$  terms are nearly equal to the hydrogen terms but the  $S$  and  $P$  terms are much larger. It is supposed that the  $S$  and  $P$  electron orbits penetrate into the group of 10 electrons around the nucleus and so are acted on by the strong field of the nucleus, which increases the energy of the orbits. All these terms are for a neutral sodium atom, and correspond to the different possible orbits of the outer electron. The first term of the principal series according to this diagram is due to a transition from the orbit with principal number 3 and second number 1 to the orbit with principal number 3 and the second number 2. This series is represented by  $P(n) = 1S - nP$ , so that in this case the principal quantum number is equal to  $n + 1$  since  $n = 2$  for the first line of the principal series.

The possible transitions, with emission of light, on the diagram are

from any dot to another dot which lies to the left on the next line above or the next line below, and so corresponds to less energy, because the energy is proportional to a constant minus the term values.

If  $n$  is the principal and  $n_1$  the second quantum number of an orbit or atomic energy level, then the level may be designated by  $n$  with a suffix  $n_1$ . Thus  $3_1$  denotes the lowest energy level of the neutral sodium atom according to the diagram.

## 8. Series Lines and Absorption.

The normal state of an atom is the state of smallest energy. An atom in a given state can only absorb light the quanta of which are equal to the quanta which the atom emits when it changes from another state of greater energy to its actual state. Thus, according to the quantum theory, sodium vapour at comparatively low temperatures at which all the atoms are in the normal  $3_1$  state should absorb light with the frequencies of the lines of the principal series. This is found to be the case. When white light is passed through sodium vapour the absorption spectrum obtained shows the principal series lines only. The same thing is true of other elements. If the frequency of the incident light is greater than that of the limit of the principal series, the outer electron may be removed right out of the atom by a quantum of the incident light. The light is absorbed in this case and the vapour is ionized. It is found that the absorption spectrum shows continuous absorption of frequencies greater than the limit of the principal series, exactly in agreement with this deduction from the quantum theory.

## 9. Doublets and Triplets.

The lines of many series are not single lines but doublets, triplets, or more complicated groups of lines. The  $P$  terms are double or triple and the different values may be indicated by a suffix thus,  $mP_i$ , where  $i = 1$  or  $2$  for a series of doublets, and  $i = 1, 2$ , or  $3$  for a series of triplets. If the  $P$  terms are double, then the  $S$  and  $D$  series lines are doublets with constant frequency differences, since  $S(n) = 2P - nS$  and  $D(n) = 2P - nD$ . The constant frequency difference is  $2P_1 - 2P_2$ . In this case the  $P$  series lines are doublets with frequency difference  $nP_1 - nP_2$ . This difference diminishes to zero as  $n$  increases, since both  $nP_1$  and  $nP_2$  tend to zero.

On the quantum theory the double nature of a term is taken to indicate a third type of quantizing requiring a third quantum number  $n_2$  to fix the state of an atom. This number is called the *inner quantum number*.

The lines of the principal series of the alkali metals are doublets. The well-known  $D$  lines of sodium are the first doublet in the sodium  $P$  series. Millikan and Bowen have observed the first doublet of the principal series of the atoms Li, Be<sub>I</sub>, B<sub>II</sub>, C<sub>III</sub>, N<sub>IV</sub>, and O<sub>V</sub>, which all contain three electrons. If  $\Delta\nu$  denotes the frequency difference of one of these doublets then they find that  $\sqrt[3]{\Delta\nu/0.365}$  is nearly equal to  $Z - 2$  in each case. For example, for lithium  $Z - 2 = 1$  and  $\sqrt[3]{\Delta\nu/0.365} = 0.981$ , and for oxygen  $Z - 2 = 6$  and  $\sqrt[3]{\Delta\nu/0.365} = 6.184$ . According to Sommerfeld's theory of the fine structure of the spectral lines of H, He<sub>I</sub>, Li<sub>II</sub> discussed in the chapter on the quantum theory,  $\Delta\nu$  for the  $H_\alpha$  line,

the first line in the Balmer series, is 0.365, and for other similar doublets should be proportional to the fourth power of the nuclear charge  $Ze$ . In the case of the atoms Li, Be, B<sub>II</sub>, C<sub>III</sub>, N<sub>IV</sub>, and O<sub>V</sub> the charge on the nucleus and group of two electrons near the nucleus is  $(Z - 2)e$ , so that on Sommerfeld's theory we should expect  $\Delta\nu$  to be nearly equal to  $0.365(Z - 2)^4$ , which is just what Millikan and Bowen find. Similar doublets with separations given by about  $0.365(Z - s)^4$ , where  $s$  is about 3.5, have been found in the X-ray spectra of practically all the elements.

## 10. Band Spectra. Quantum Theory.

The spectra of many compounds and of diatomic elements contain groups of lines which are called bands. In these bands the lines get closer together from one side of the band to the other, so that one side of the band has a well-defined edge called the head of the band where the lines are very close together. Such spectra frequently contain a series of such bands, all with heads on the same side. A good example of such a band spectrum is the spectrum of the light from the inner cone of a Bunsen flame, which is usually attributed to carbon monoxide. Cyanogen and hydrochloric acid also give band spectra. Many elements give so-called many-lined spectra, which are probably band spectra in which the different bands overlap and so become confused. An example of such a spectrum is the many-lined spectrum of hydrogen, which is attributed to H<sub>2</sub>. Several bands have been located in this spectrum by O. W. Richardson and others.

According to the quantum theory the bands are emitted by rotating molecules. The rotating molecules are supposed to be only able to exist in a number of definite states characterized by definite rotational energies, and to emit light when they change from one such state to another with less energy. The frequency of the light emitted is given by the usual equation  $h\nu = E_1 - E_2$ . Let  $K$  denote the moment of inertia of a molecule about its axis of rotation, and  $\theta$  the angle through which it has turned. Then according to the quantum theory we have  $\int_0^{2\pi} K\dot{\theta}d\theta = n\hbar$ , where  $n$  is an integer. Hence, since  $\dot{\theta}$  is constant,  $K\dot{\theta} = n\hbar/2\pi$ .

The kinetic energy of the molecule is therefore given by

$$E_n = \frac{1}{2}K\dot{\theta}^2 = \frac{1}{8\pi^2} \frac{\hbar^2 n^2}{K}.$$

We assume that  $n$  can only change by one unit at a time, so that when it is  $+$  it can change to  $n - 1$ , and when  $-$  to  $n + 1$ . The frequency of the light emitted is then

$$\nu = \frac{E_n - E_{n-1}}{\hbar} = \frac{\hbar}{8\pi^2 K} \{n^2 - (n-1)^2\} = \frac{\hbar}{8\pi^2 K} (2n-1),$$

when  $n$  is  $+$ , and  $\nu = -\frac{\hbar}{8\pi^2 K} (2n+1)$ , when  $n$  is  $-$ .

The frequencies emitted are therefore

$n$ .		$\nu$ .
$+4$	....	$7\nu_0$
$+3$	....	$5\nu_0$
$+2$	....	$3\nu_0$
$+1$	....	$\nu_0$
$0$	....	$-\nu_0$
$-1$	....	$\nu_0$
$-2$	....	$3\nu_0$
$-4$	....	$7\nu_0$

where  $\nu_0 = h/8\pi^2 K$ . The frequency  $-\nu_0$  for  $n=0$  indicates an absorption of energy by the molecule. The spectrum should therefore consist of lines with frequencies  $\nu_0, 3\nu_0, 5\nu_0$ , &c. Such spectra have been observed far in the infra-red, notably for water vapour, by Rubens and Bahr.

If we suppose that the molecule also vibrates with a frequency  $\nu_1$ , then according to the quantum theory its vibrational energy can only change by  $h\nu_1$ , so that the frequencies which will be emitted when the vibrational and rotational energies both change will be given by

$$\nu = \nu_1 + \frac{h}{8\pi^2 K} \{n^2 - (n \pm 1)^2\},$$

or

$$\nu = \nu_1 + \nu_0 (\pm 2n - 1).$$

In this case, when  $\nu_1$  is greater than  $\nu_0$ , as is usually the case,  $n$  can change to  $n+1$  or to  $n-1$ , with emission of light. The frequencies emitted are

$n$ .		$\nu$ .	
4	....	$\nu_1 + 7\nu_0$	$\nu_1 - 9\nu_0$
3	....	$\nu_1 + 5\nu_0$	$\nu_1 - 7\nu_0$
2	....	$\nu_1 + 3\nu_0$	$\nu_1 - 5\nu_0$
1	....	$\nu_1 + \nu_0$	$\nu_1 - 3\nu_0$
0	....	$\nu_1 - \nu_0$	$\nu_1 - \nu_0$

Thus a band is obtained with frequencies  $\nu_1 + \nu_0, \nu_1 - \nu_0, \nu_1 + 3\nu_0, \nu_1 - 3\nu_0$ , and so on, but not with frequency  $\nu_1$ . Such bands are observed in the infra-red spectra of HCl, HBr, and other diatomic molecules. The line  $\nu_1 + \nu_0$  is usually absent.

So far we have supposed that the moment of inertia of the molecule is a constant independent of its rotational and vibrational energies, and that the internal energy of the molecule does not change. In the case of bands in the visible or ultra-violet regions it is necessary to suppose that the internal energy of the molecule changes as well as the rotational and vibrational energies. The change in the internal energy, however, may be supposed to involve a change in the moment of inertia and so of  $\nu_0$ . The frequencies emitted are therefore given by

$$\nu = \nu_1 + \nu_2 + \nu_0 n^2 - \nu_0' (n \pm 1)^2,$$

where  $\nu_0'$  is the new value of  $\nu_0$ , and  $\nu_2$  corresponds to the change in the internal energy.

This gives

$$\nu = A \pm Bn + Cn^2,$$

where

$$A = \nu_1 + \nu_2 - \nu_0', \quad B = 2\nu_0', \quad C = \nu_0 - \nu_0'.$$

It is found that many bands can be represented very well by this expression. The frequencies given by  $\nu_- = A - Bn + Cn^2$  are said to form the negative branch of the band, and those given by  $\nu_+ = A + Bn + Cn^2$  the positive branch.

The head of the band is at the minimum value of  $\nu$  on the negative branch, and is given by  $\frac{d\nu_-}{dn} = -B + 2Cn = 0$ , or  $n = \frac{B}{2C}$ , so that

$$\nu_- = A - B \frac{B}{2C} + C \frac{B^2}{4C^2} = A - \frac{B^2}{4C}.$$

If  $B/2C$  is not equal to an integer then the head is at the value of  $\nu$  given by

the integer nearest to  $B/2C$ . The relation between  $\nu$  and  $n$  can be made very clear graphically, as shown in fig. 2.

The curves represent  $n$  as a function of  $\nu$ , and horizontal lines are drawn to

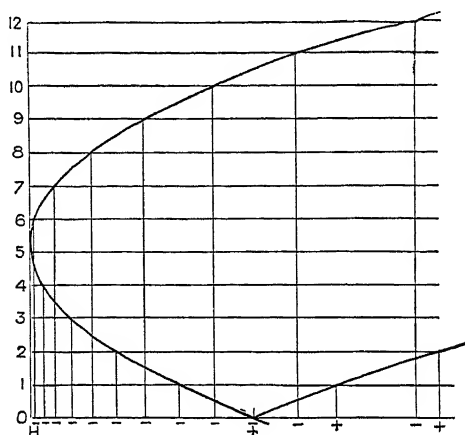


Fig. 2

represent the positive integral values of  $n$ . The values of  $\nu$  in the band are then indicated by the intersections of the horizontal lines and the curves. The head of the band is at H.

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## CHAPTER IX

# Cathode Rays, $\beta$ -Rays, and $\alpha$ -Rays

### CATHODE RAYS

#### 1. Crookes' Experiments.

Cathode rays were discovered by Plücker in 1859; he observed that when an electric discharge was passed between two electrodes through a tube and the gas pressure was reduced to a sufficiently small value, the glass walls of the tube near the cathode emitted a greenish-coloured light. This light could be moved about by bringing a magnet near the tube, and appeared to be produced by something coming from the cathode and striking the glass. Cathode rays were later investigated by Hittorf and Goldstein and in 1879 by Crookes, whose beautiful experiments made clear many of the properties of the rays in a striking way.

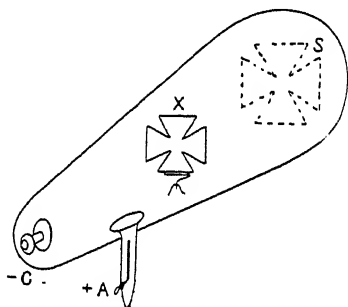


Fig. 1

As the pressure in a discharge tube is reduced the distance between the negative glow and the cathode increases, and when the Crookes dark space reaches the walls of the tube the green light observed by Plücker becomes very bright.

If a solid body is put up between the cathode and the walls of the tube a sharply defined shadow of it appears in the green light emitted by the walls. This was clearly shown by Crookes with the famous "Crookes tube" shown in fig. 1. The glass tube is conical in shape with a nearly flat end at S. At C a small aluminium disc supported by a wire sealed through the glass serves as the cathode. The anode is a wire A in a side tube. At X a piece of mica or aluminium sheet in the form of a cross is supported on a hinge so that by tilting the tube it can be put in either a horizontal or vertical position. The gas pressure in the tube is reduced to about 0.01 mm., so that the Crookes dark space fills the tube and the greenish light is emitted by the flat

end when the cathode and anode are connected to an induction coil. When the cross is up it throws a sharp shadow on the end of the tube, showing clearly that something is emitted by the cathode which is stopped by the cross. Also the shadow is sharp although the source is of considerable size, showing that the rays are not emitted in all directions from every point on the cathode but in only one direction. It is found that the rays are emitted in a direction perpendicular to the surface of the cathode close to its surface. Thus Crookes showed that by using a concave cathode the rays could be focused on to a small area, and if a piece of platinum foil was put up at the focus the foil became very hot and could even be melted. The cathode rays are deflected by a magnetic field as we should expect negatively charged particles to be deflected. A Crookes tube for showing this is shown in fig. 2.

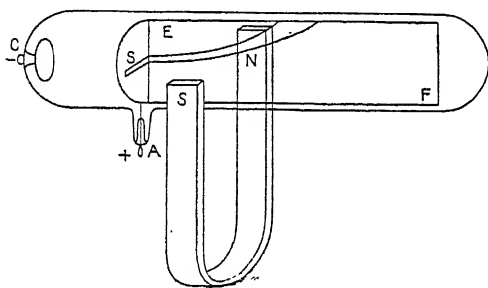


Fig. 2

The cathode is a disc at C. Parallel to it a few centimetres away is a metal sheet, with a slit S cut in it, which forms the anode. The narrow beam of cathode rays which pass through the slit falls on a slightly inclined screen EF coated with calcium sulphide, or some

other substance, which fluoresces brightly when struck by the rays. A bright streak of light appears on the screen which can be deflected by a magnetic field perpendicular to the screen. In the figure a magnet is shown with its S pole in front and its N pole behind the tube, deflecting the rays upwards as shown.

## 2. The Wehnelt Cathode. Magnetic and Electric Deflection.

It was found by Wehnelt that calcium and barium oxides emit cathode rays very freely when heated to a red heat and negatively charged. By means of a cathode consisting of a strip of platinum foil, which can be heated by a current, having a small area on it coated with calcium or barium oxide an intense narrow beam of cathode rays can be obtained. If the gas pressure is not too low the narrow beam causes the gas to emit light so that its path through the tube can be easily seen in a dark room and the deflection of such a beam by a magnetic field observed. A tube with a Wehnelt cathode is shown in fig. 3. The anode A is a small aluminium disc and the cathode C a strip of platinum foil about 2 mm. wide supported by two wires E and F. A small patch of oxide is on the strip near its middle. E and F are

connected to a battery and the strip heated to a dull red heat. A potential difference of a few hundred volts between C and A is then sufficient to produce an intense beam of cathode rays from the oxide patch which comes out perpendicular to the surface of the foil and diverges slightly. By means of a magnet the beam of rays can be deflected along a curve as shown in the figure.

The potential difference required to produce a discharge in a Crookes tube, in which the dark space fills the tube, is usually of the order of 10,000 volts or more, which is much greater than is necessary with a Wehnelt cathode. The magnetic deflection of cathode rays increases as the potential difference used to produce them decreases, so that the rays from a Wehnelt cathode using a few hundred volts are much more easily deflected than the rays in a Crookes tube. Cathode rays are deflected by an electric field like negatively charged particles. A tube used by J. J. Thomson to show this is shown in fig. 4. The

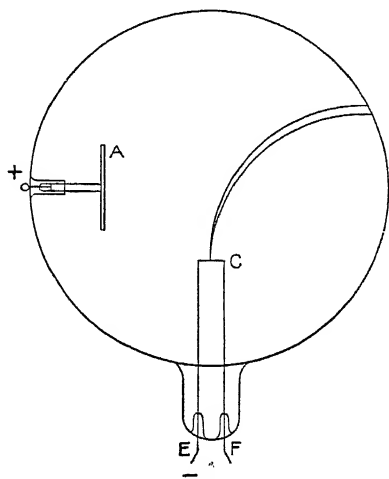


Fig. 3

cathode is a small aluminium disc at C. The anode is a metal disc A having a small hole in it through which a narrow beam of the rays from C passes. The beam is further limited by a second disc B with a small hole in it, and then passes between two parallel metal plates D and E to the end of the tube near P, where it produces a luminous spot on the glass. If D and E are connected together the rays pass along the axis of the tube, but if a potential difference is maintained between them by means of a battery the rays are deflected towards the positively charged plate.

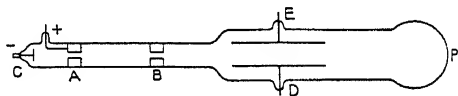


Fig. 4

That the cathode rays carry a negative charge was

first shown directly by Perrin in 1895. He passed a beam of the rays into a metal cylinder and found that the cylinder received a negative charge. Hertz noticed that cathode rays can pass through very thin metal foil, and Lenard showed that they can be passed through a thin metal window in a discharge tube into the air outside or into another tube containing a gas at any desired pressure.

### 3. Cathode Rays are Negatively Charged Particles. Charge and Mass.

The experimental results described above show that cathode rays are negatively charged particles moving with a high velocity, and for a time it was generally believed that they were negatively charged atoms of the gas in the discharge tube or of the metal of the cathode. About 1897, however, several physicists succeeded in measuring the ratio of the charge  $e$  to the mass  $m$  of the cathode ray particles and found this ratio to have a surprisingly high value, about  $10^7$  electromagnetic units per gramme. It was also shown by Lenard that they have considerable penetrating power and can pass through several millimetres of air at atmospheric pressure.

Before 1897 the highest ratio of charge to mass known was that for the positively charged hydrogen ion in solution for which  $e/m$  is about  $10^4$  E.M. units per gramme. The charge on a univalent ion in solutions was believed to be a sort of atomic unit of electricity because ionic charges in solutions are always multiples of it, so that physicists were inclined to expect that gaseous ions would be found to carry the same charges as ions in solutions. The high value of  $e/m$  for cathode rays might have been explained by supposing  $e$  to be about 1000 times the charge on one hydrogen ion in solutions and  $m$  equal to the mass of one hydrogen atom. J. J. Thomson, who was one of the first to measure  $e/m$  for cathode rays, pointed out that the cathode rays could not have the high penetrating power observed by Lenard if they were of atomic dimensions, and suggested that the high value of  $e/m$  was due to  $m$  being about 1000 times smaller than the mass of one hydrogen atom, the charge  $e$  being the same as that on one univalent ion in solutions. J. J. Thomson supported this revolutionary theory by many ingenious experiments, and it is now universally accepted. Other physicists, notably Wiechert and Kaufmann, also measured  $e/m$  for cathode rays at the same time as J. J. Thomson, and Wiechert suggested the same explanation of its high value, but to J. J. Thomson as the principal advocate of the new theory belongs the greater share of the credit for it.

The cathode ray particles or electrons, as they are now called, can be obtained from all kinds of matter, and are one of the two constituents of which atoms appear to be composed. The discovery of particles 1000 times lighter than hydrogen atoms, obtainable from all kinds of matter, may be said to mark the beginning of modern physics as distinguished from the classical physics of Faraday, Clerk-Maxwell, and Helmholtz.

#### 4. Ratio of Charge to Mass. Measurements of Kaufmann and J. J. Thomson.

The methods used to measure  $e/m$  for cathode rays will now be considered. The apparatus used by Kaufmann is shown in fig. 5. The cathode C and anode A were connected to a Wimshurst machine, and the potential difference  $V$  between them was measured with an electrostatic voltmeter. The anode was a thin straight wire, and the cathode rays produced a shadow of it on the plate P which closed the lower end of the tube. A nearly uniform magnetic field perpendicular to the rays and parallel to the anode wire was produced by a large coil SS' which surrounded the part of the tube below the anode as shown. The magnetic field deflected the rays and the deflection of the shadow of the anode on the plate P was measured. The velocity  $v$  of an electron starting from the cathode is given by the equation

$$Ve = \frac{1}{2}mv^2,$$

where  $e$  is the charge and  $m$  the mass of the electron. In the magnetic field ( $H$ ) the electrons move along a circular path of radius  $r$  given by  $mv^2/r = Hev$ . The radius  $r$  was calculated from the observed deflection and the dimensions of the apparatus. These equations give  $e/m = 2V/H^2r^2$ , and  $v = 2V/Hr$ . The potential difference  $V$  was varied from about 3000 to 10,000 volts by reducing the gas pressure in the tube from 0.07 mm. to 0.03 mm. The final result obtained was  $\frac{e}{m} = 1.77 \times 10^7$  E.M. units per gramme.

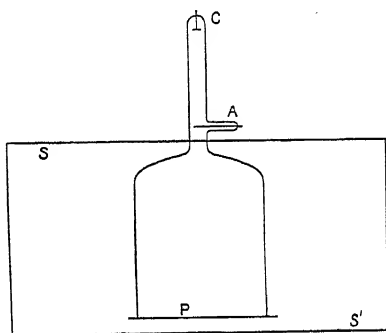


Fig. 5

A possible objection to this method is that some of the electrons may be liberated in the gas between the cathode and anode and so not fall through the whole of the potential difference  $V$ . Such electrons, however, would not produce a sharp shadow on the screen because they would be deflected through different distances depending on their velocity. The sharp shadow which was obtained must have been due to electrons all having the same velocity, so that they must have started at the surface of the cathode as Kaufmann assumed. In discharges at low pressures the cathode rays are mainly produced by the impact of positive ions on the cathode.

J. J. Thomson, using a tube like that shown in fig. 4, measured the deflection of the rays by a magnetic field and then balanced the

magnetic deflection by the electric field between the plates D and E. The magnetic field was perpendicular to the rays and parallel to the surfaces of the plates D and E, and extended over the space between these plates. In this case  $Fe = Hev$ , where  $F$  is the electric field strength, so that  $v = F/H$ . For the magnetic deflection  $mv^2/r = Hev$ , so that  $e/m = F/H^2r$ . In this way J. J. Thomson found  $e/m$  to be about  $10^7$ , and  $v$  to be about  $3 \times 10^9$  cm. per second when the gas pressure was 0.01 mm.

J. J. Thomson also measured  $e/m$  by another method. A beam of the rays was received in a small metal cylinder, and the rate of rise of temperature of this cylinder was determined with a thermocouple. The charge received by the cylinder was measured by connecting it to a galvanometer. In this way the energy  $W$  and charge  $Q$  of the rays were found. We have then  $W/Q = \frac{1}{2}mv^2/e$ , so that if  $v$  is found from the magnetic and electric deflections or otherwise,  $e/m$  can be calculated. In this way J. J. Thomson found  $e/m$  to be about  $10^7$ , as in his previous experiments.

The value of  $e/m$  for cathode rays has, since 1897, been carefully measured by many physicists. The value now accepted for slowly moving electrons is  $1.769 \times 10^7$  E.M. units per gramme. It was found that  $e/m$  for the cathode rays is the same for discharges in different gases and with cathodes made of different metals. It is also the same for the cathode rays emitted by hot bodies or set free by ultra-violet light, and for the  $\beta$ -rays or electrons emitted by radioactive bodies. It appears that electrons of identical properties can be obtained from any kind of matter; they must therefore be one of the constituents of the chemical atoms.

## 5. Cathode Rays and Ionization of Gases.

When cathode rays are passed through a gas they produce ions by collisions with the gas molecules. The ionization produced by electrons moving through a gas under the action of a uniform electric field is discussed in the chapter on the motion of electrons in gases. When cathode rays are passed into a gas the velocity with which they move is gradually diminished by collisions with the gas molecules. High velocity rays travel in straight lines through gases at low pressures with little loss of energy, but slower rays are deflected by the molecules and describe paths of irregular shape until they are stopped. The number of pairs of ions produced by cathode rays per centimetre of path has been determined by Durack and by Glasson. Durack found that cathode rays produced 0.4 pair of ions and that  $\beta$ -rays from radium produced 0.17 pair of ions per centimetre in air at 1 mm. pressure. The velocity of the cathode rays was probably about  $4 \times 10^9$  cm. per second and that of the  $\beta$ -rays about  $2 \times 10^{10}$  to  $2.8 \times 10^{10}$  cm. per second.

Glasson obtained cathode rays of known velocity by deflecting them by a uniform magnetic field and obtained the following results:

Velocity in Cm./Sec.		Pairs of Ions produced per Cm. in Air at 1 Mm. Pressure.
$4.08 \times 10^9$	....	2.01
$4.76 \times 10^9$	....	1.53
$5.44 \times 10^9$	....	1.26
$6.12 \times 10^9$	.. .	0.99

It appears that the ionization produced diminishes as the velocity increases. The number of molecules struck by an electron in going 1 cm. through air at 1 mm. pressure must, according to the kinetic theory of gases, be about 180. For more slowly moving electrons Townsend found the maximum number of pairs of ions formed to be 14.6 per centimetre in air at 1 mm. pressure.

## 6. Absorption of Cathode Rays.

The absorption of cathode rays by thin metal sheets and by gases was investigated by Lenard in 1895. The cathode rays in a discharge tube were passed, through a small window of aluminium foil in the wall of the tube, into another tube. A screen of barium platinocyanide was put up in the second tube opposite the window so that the cathode rays fell on it and caused it to phosphoresce. The intensity of the rays at the screen was estimated by the brightness of the luminosity of the screen. The absorption of the rays by thin sheets of metal and by gases at different pressures was determined by interposing them between the window and the screen and measuring the diminution of the luminosity. It was found that the intensity of the rays transmitted was nearly proportional to  $\epsilon^{-\lambda t}$  where  $\lambda$  is a constant and  $t$  the thickness of the absorbing substance. The constant  $\lambda$  depends on the nature of the absorbing substance and on the velocity of the rays. It was found that for rays produced by a potential difference of 30,000 volts the constant  $\lambda$  was nearly proportional to the density  $d$  of the absorbing layer. The following table gives some of Lenard's results.

Substance.	$\lambda$ cm. <sup>-1</sup> .	$d$ gm./cm. <sup>3</sup> .	$\lambda/d$ .
Hydrogen at 3 mm. pressure ..	0.00149	$3.5 \times 10^{-7}$	4,040
Hydrogen at 760 mm. pressure ...	0.476	$8.5 \times 10^{-5}$	5,610
Air at 760 mm. pressure ..	3.42	$1.2 \times 10^{-3}$	2,780
Paper .. .. .	2,690	1.3	2,070
Mica .. .. .	7,250	2.8	2,590
Gold .. .. .	55,600	19.3	2,880

Thus it appears that for a range of  $d$  from  $3.6 \times 10^{-7}$  to 19.3 the quotient  $\lambda/d$  only varies by a factor of less than 3. This extremely interesting result was obtained by Lenard in 1895. If  $M$  denotes the mass per unit area of the absorbing layer then  $M = dt$ , so that  $\lambda t = \lambda M/d$ . Since  $\lambda/d$  is nearly the same for all substances this shows that the absorption depends almost entirely on the mass of the screen and not on the kind of matter of which it is made. According to the modern theory of atoms each atom consists of a positively charged nucleus sur-

rounded by a number of electrons. The number of electrons is roughly half the atomic weight, so that the number of electrons per cubic centimetre in any kind of matter is roughly proportional to the density of the matter. In the case of heavy atoms like gold the number of electrons is much greater than the number of nuclei, but in the case of hydrogen there is only one electron to each nucleus. Thus the sum of the numbers of electrons and nuclei per unit mass of matter for hydrogen is about double its value for the heavier elements. The absorption of the cathode rays may be attributed to collisions with the electrons and nuclei, so that we should expect the ratio  $\lambda/d$  for hydrogen to be about double its value for the heavier elements, as was found by Lenard to be the case. The absorption of cathode rays is nearly the same for different kinds of matter because all kinds of matter consist of electrons and nuclei. The electrons and nuclei only occupy a very minute fraction of the volume of the matter so that cathode rays are able to pass through. We should expect any kind of atom to pass through matter as easily as cathode rays, but we cannot obtain atoms moving with the high velocity of cathode rays.

W. Wilson measured the velocity  $V$  of cathode rays before and after passing them through thin sheets of metal. The velocity was found from the radius of the circle described in a magnetic field of known strength. He found that

$$V_0^4 - V^4 = kt,$$

where  $V_0$  is the velocity before entering the sheet,  $V$  that after passing through,  $t$  the thickness of the sheet, and  $k$  a constant. The rays are deflected in the sheet by collisions with the atoms, so that the distance they actually travel in the sheet may be considerably greater than  $t$ . Since the energy of the rays is nearly proportional to  $V^2$  and the number of collisions to  $t$ , this indicates that the energy lost per collision is inversely as the energy of the rays.

### $\beta$ -RAYS

#### 7. Kaufmann's Experiments.

The  $\beta$ -rays which are emitted by many radioactive substances are found to be of precisely the same nature as cathode rays. They carry a negative charge and are deflected by electric and magnetic fields. The velocities of  $\beta$ -rays range up almost to the velocity of light.

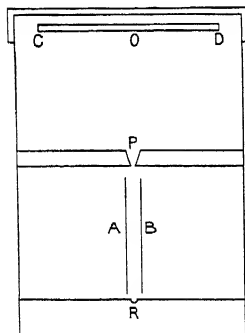


Fig. 6

The ratio  $e/m$  for  $\beta$ -rays was investigated accurately by Kaufmann and later by Bucherer. Kaufmann's apparatus is shown in fig. 6. A particle of radium bromide was placed at  $R$  in a depression in a thick metal plate. The  $\beta$ -rays from the radium passed between two parallel metal plates  $A$  and  $B$  to a small hole  $P$  in a thick metal sheet. From  $P$  the narrow beam of rays passed on to a photographic plate  $CD$ . A potential difference was maintained between  $A$  and  $B$  by means of a battery, and the whole apparatus was placed in a uniform magnetic

field parallel to the direction of the electric field between  $A$  and  $B$ . The magnetic field deflects the rays so that they move along circular

paths, in planes perpendicular to the field, of radius  $r$  given by  $mv^2/r = Hev$ . The circular path of the rays is shown in fig. 7. The radius is at  $R$  and the rays pass through the small hole at  $P$  and fall on the photographic plate at  $Q$  so that  $R$ ,  $P$ , and  $Q$  lie on the circle. Let the magnetic deflection  $OQ = y$ ,  $RO = a$ , and  $RP = b$ . Then we have

$$r^2 = \frac{b^2}{4} + \left( \frac{a^2 - ab + y^2}{2y} \right)^2,$$

so that  $r$  can be calculated when  $y$ ,  $a$ , and  $b$  have been measured.

The electric deflection, which will be denoted by  $z$ , is perpendicular to  $y$  on the photographic plate. To calculate  $z$  approximately we may assume the electric field uniform, and equal to  $F$  from  $R$  to  $P$ , and to zero from  $P$  to the plate. The path of the rays in the plane of the electric field is shown in fig. 8. It is parabolic from  $R$  to  $P$ , and straight from  $P$  to  $Q'$ . The acceleration of the electrons due to the electric field is  $Fe/m$ , so that approximately

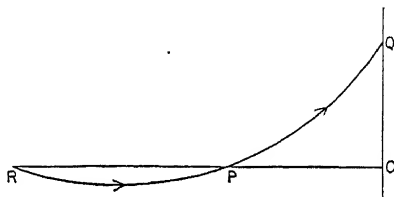


Fig. 7

$\frac{1}{2} \left( \frac{Fe}{m} \right) \left( \frac{b}{v} \right)^2 - v' \frac{b}{v} = 0$ , where  $v'$  is the initial transverse velocity component of the rays which get to  $P$ . Hence  $v' = \frac{1}{2} \left( \frac{Fe}{m} \right) \frac{b}{v}$ . The transverse velocity at  $P$  due to

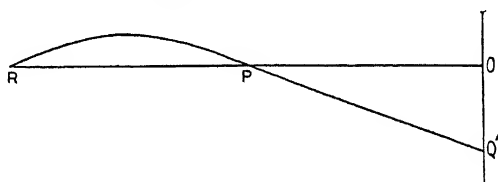


Fig. 8

the electric field is equal to  $\frac{Fe b}{m v}$ , so that the resultant transverse velocity at  $P$  is  $\frac{1}{2} \frac{Fe b}{m v}$ . Hence

$$z = (a - b) \frac{\frac{1}{2} \frac{Fe b}{m v}}{v},$$

or

$$z = (a - b) \frac{b}{2} \frac{Fe}{mv^2}.$$

This equation and  $mv/r = He$  enable  $v$  and  $e/m$  to be calculated. The  $\beta$ -rays emitted by radium bromide have velocities varying from  $10^{10}$  cm. per second to almost  $3 \times 10^{10}$  cm. per second. Kaufmann thus obtained a curve on his photographic plates the co-ordinates of points on which were  $y$  and  $z$ . Each point on the curve corresponds to rays having a definite velocity, so that by measuring  $y$  and  $z$  for different points on the curve  $e/m$  could be found for rays having different velocities. Kaufmann found that  $e/m$  decreases as  $v$  increases. The variation observed agrees nearly with the equation

$$\frac{e}{m} = \frac{e}{m_0} \sqrt{1 - v^2/c^2},$$

where  $c$  is the velocity of light and  $e/m_0$  is the value of  $e/m$  when  $v/c$  is small.

### 8. Bucherer's Experiments. Mass and Velocity.

The apparatus used by Bucherer is shown in fig. 9. AB and CD are two parallel circular metal discs only a fraction of a millimetre apart. A small particle of radium, R, was put between them at their centres. The  $\beta$ -rays from the radium moved out from R radially in all directions between the discs and fell on a cylindrical

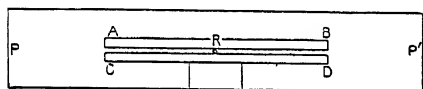


Fig. 9

photographic film PP' which was concentric with the discs. The discs and film were in a metal box in which a good vacuum was maintained. An electric field  $F$  was produced between the discs by connecting them to a battery, and the whole apparatus was put in a uniform magnetic field  $H$  parallel to the plane of the discs. Since the discs were so near together the electrons from the radium could not get to the photographic film unless the force on them due to the electric field was equal and opposite to that due to the magnetic field. We have therefore, for rays going along a radius making an angle  $\theta$  with the magnetic field,

$$Fe = Hev \sin \theta, \text{ or } v = F/H \sin \theta.$$

Bucherer made  $F/H$  equal to  $\frac{1}{2}c$  in most of his experiments, so that  $v/c = 1/2 \sin \theta$ . If  $v = c$  this gives  $\sin \theta = \frac{1}{2}$ , so that  $\theta = 30^\circ$  or  $150^\circ$ . Now  $v$  is never greater than  $c$ , so that no rays should get to the film except between  $\theta = 30^\circ$  and  $\theta = 150^\circ$ . Bucherer found that this was the case. When  $\theta = \pi/2$  then  $v = \frac{1}{2}c$ , which is therefore the smallest velocity of the rays which can get to the film when  $F/H$  is equal to  $\frac{1}{2}c$ . The rays which get out from between the discs are deflected by the magnetic field and so give a curve on the photographic film. The

deflection is approximately equal to  $\frac{1}{2}a^2He \sin\theta/mv$ , where  $a$  is the difference between the radius of the film and that of the discs, for the magnetic field gives the electrons an acceleration  $Hev \sin\theta/m$  which acts for a time  $a/v$ .

In this way Bucherer found  $v$  and  $e/m$  for several values of  $v$ . The following table gives his results:

$v/c$ .		$\frac{e \times 10^{-7}}{m \sqrt{1 - v^2/c^2}}$ .
0.3173	....	1.752
0.3787	....	1.761
0.4281	....	1.760
0.5154	....	1.763
0.6870	....	1.767

The second column gives the observed values of  $e/m$  multiplied by  $10^{-7}$  and divided by  $\sqrt{1 - v^2/c^2}$ . It appears that this quantity is practically constant and equal to  $10^{-7} \times e/m$  for cathode rays, which is 1.769. If we assume that the charge  $e$  is independent of  $v$  then these experiments show that

$$m = \frac{m_0}{\sqrt{1 - v^2/c^2}},$$

where  $m_0$  is the mass when  $v = 0$ .

This variation of the mass with the velocity is shown in the chapters on the electron theory and on relativity to be due to energy possessing mass equal to the energy divided by the square of the velocity of light, so that when the kinetic energy of a particle of any kind is increased then its mass is also increased. At one time it was supposed that the observed variation of the mass of electrons with their velocity gave information as to the nature of these particles, but it is now clear that if energy possesses momentum equal to  $Ev/c^2$ , where  $E$  is the energy,  $v$  its velocity, and  $c$  the velocity of light, then the mass of any particle varies with its velocity so that

$$m = \frac{m_0}{\sqrt{1 - v^2/c^2}}.$$

The observed variation of the mass of electrons with their velocity is therefore presumably the same as for all other bodies and gives no information about their constitution or the distribution of the charge in them. So far as we know, electrons behave like particles of mass  $m$  and charge  $e$ , so that the equation of motion of an electron is

$$\frac{d}{dt}(mv) = F,$$

where  $F$  is the resultant force on the electron and  $mv$  its momentum.

### 9. Scattering of $\beta$ -rays by Matter. Mathematical Theory.

When  $\beta$ -rays are passed through thin sheets of matter they are scattered like cathode rays, but owing to their higher velocities the scattering is less. If a narrow parallel beam of  $\beta$ -rays is allowed to fall normally on a thin metal sheet, of thickness  $t$ , then the rays which emerge on the other side form a diverging beam the intensity in which is greatest along the normal to the sheet and falls off gradually as the inclination to the normal is increased. The angle between the emerging  $\beta$ -rays and the normal, or the deviation from their original direction, is called the scattering angle and will be denoted by  $\varphi$ . In passing through the sheet any particular  $\beta$ -ray passes through a large number of atoms, so that the angle  $\varphi$  through which it is finally deviated is the resultant of a large number of small deviations.

Let the initial direction of the beam be horizontal and let the deviation of a  $\beta$ -ray in the horizontal plane be  $x$  and in the vertical plane  $y$ , so that when  $\varphi$  is small we have  $\varphi^2 = x^2 + y^2$ . Let the number of rays for which  $x$  is between  $x$  and  $x + dx$  be

$$n_0 f(x) dx,$$

where  $n_0$  is the total number of  $\beta$ -rays considered, and  $f(x)$  denotes a function of  $x$  only. In the same way let  $n_0 f(y) dy$  denote the number for which  $y$  is between  $y$  and  $y + dy$ . Then, since it is clear that the distribution as regards  $x$  is independent of that as regards  $y$ , the number for which  $x$  is between  $x$  and  $x + dx$  and also  $y$  between  $y$  and  $y + dy$  must be

$$n_0 f(x) f(y) dx dy.$$

But this must be a function of  $x^2 + y^2$  since the distribution must clearly be symmetrical about the origin at  $x = 0$  and  $y = 0$ . Hence

$$n_0 f(x) f(y) dx dy = n_0 \psi(x^2 + y^2) dx dy,$$

where  $\psi(x^2 + y^2)$  denotes some function of  $x^2 + y^2$ . Hence

$$f(x) f(y) = \psi(x^2 + y^2).$$

Differentiating this with respect to  $x$  and dividing one side by  $f(x)f(y)$  and the other by  $\psi(x^2 + y^2)$  we get

$$\frac{f'(x)}{f(x)} = 2x \frac{\psi'(x^2 + y^2)}{\psi(x^2 + y^2)}.$$

In the same way

$$\frac{f'(y)}{f(y)} = 2y \frac{\psi'(x^2 + y^2)}{\psi(x^2 + y^2)},$$

so that

$$\frac{1}{2x} \frac{f'(x)}{f(x)} = \frac{1}{2y} \frac{f'(y)}{f(y)}.$$

Hence both these quantities must be equal to a constant, say  $-\alpha$ , so that

$$\frac{f'(x)}{f(x)} = -2\alpha x,$$

which gives

$$\log f(x) = -\alpha x^2 + \text{constant},$$

or

$$f(x) = A e^{-\alpha x^2},$$

where  $A$  is another constant. To determine the constants we have

$$n_0 \int_{-\infty}^{+\infty} f(x) dx = n_0,$$

and, if  $\bar{x}^2$  denotes the average value of  $x^2$ ,

$$n_0 \int_{-\infty}^{+\infty} x^2 f(x) dx = n_0 \bar{x}^2.$$

These equations give  $A \sqrt{\frac{\pi}{\alpha}} = 1$  and  $\frac{A}{2\alpha} \sqrt{\frac{\pi}{\alpha}} = \bar{x}^2$ , so that  $\alpha = \frac{1}{2\bar{x}^2}$ . Hence

$$f(x) = A e^{-x^2/2\bar{x}^2}.$$

and in the same way

$$f(y) = A e^{-y^2/2\bar{y}^2}.$$

Also, if  $\bar{\varphi}^2$  denotes the mean value of  $\varphi^2$  then, since  $\varphi^2 = x^2 + y^2$ , we have  $\bar{\varphi}^2 = \bar{x}^2 + \bar{y}^2 = 2\bar{x}^2$ , for  $\bar{x}^2$  must be equal to  $\bar{y}^2$ . Hence

$$\psi(\varphi^2) = f(x)f(y) = A^2 e^{-\frac{x^2+y^2}{2\bar{x}^2}} = A^2 e^{-\varphi^2/\bar{\varphi}^2}.$$

The number of  $\beta$ -rays for which  $\varphi$  is between  $\varphi$  and  $\varphi + d\varphi$  is therefore

$$n_0 A^2 e^{-\varphi^2/\bar{\varphi}^2} 2\pi\varphi d\varphi,$$

or

$$\pi n_0 A^2 e^{-\varphi^2/\bar{\varphi}^2} d(\varphi^2).$$

The number for which  $\varphi$  is less than a given value  $\varphi$  is therefore

$$\pi n_0 A^2 \int_0^{\varphi} e^{-\varphi^2/\bar{\varphi}^2} d(\varphi^2) = \pi n_0 A^2 \bar{\varphi}^2 (1 - e^{-\varphi^2/\bar{\varphi}^2}).$$

Hence, if  $n_0$  denotes the number of rays passing through the sheet and  $n$  the number for which the deviation is less than  $\varphi$ , we have

$$n/n_0 = 1 - e^{-\varphi^2/\bar{\varphi}^2}.$$

The deviation  $\varphi$  is the resultant of the deviations due to individual atoms in the sheet. Let  $x_1$  denote the horizontal deviation due to a single atom and  $y_1$  the vertical deviation. Then we have

$$\begin{aligned} x &= \Sigma x_1, \\ y &= \Sigma y_1, \end{aligned}$$

where  $x^2 + y^2 = \varphi^2$ , and  $\Sigma x_1$  denotes the sum of all the horizontal deviations of an electron while passing through the sheet. Hence

$$x^2 = (\Sigma x_1)^2 = \Sigma x_1^2,$$

because the products of two different  $x_1$ 's are as likely to be positive as to be negative, and so their sum must be zero. In the same way

$$y^2 = \Sigma y_1^2,$$

so that

$$\varphi^2 = \Sigma (x_1^2 + y_1^2) = \Sigma \varphi_1^2,$$

where  $\varphi_1$  is the deviation due to one atom. The mean value of  $\varphi^2$  is therefore given by

$$\bar{\varphi}^2 = N \bar{\varphi}_1^2,$$

where  $N$  is the number of atoms which a  $\beta$ -ray passes through in the sheet and  $\bar{\varphi}_1^2$  is the mean of the squares of the atomic deviations. Hence we have

$$n/n_0 = 1 - e^{-\varphi^2/N\bar{\varphi}_1^2}.$$

The number  $N$  will be approximately given by  $N = \pi \mathcal{N} R^2 t$ , where  $\mathcal{N}$  is the number of atoms per cubic centimetre in the sheet and  $R$  the radius of an atom, for this is the number of atoms in a cylinder of radius  $R$  and length  $t$  which a  $\beta$ -ray going straight through the sheet would pass through. Hence

$$n/n_0 = 1 - \varepsilon - \phi^2 / (\pi \mathcal{N} R^2 \phi_1^2 t).$$

### 10. Experiments on Scattering of $\beta$ -rays.

An apparatus used by Crowther to investigate the scattering of  $\beta$ -rays is shown in fig. 10. The radioactive body was placed at R in a vertical hole in a lead block.

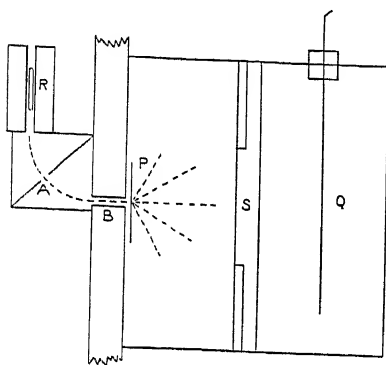


Fig. 10

The  $\beta$ -rays from R passed out downwards and were deviated sideways by means of a magnetic field perpendicular to the plane of the paper, so that they traversed a circular path RAB, the beam being limited by diaphragms so that only rays having a definite velocity got to P. At P the rays fell on a thin sheet of metal and emerged from it as a diverging beam. The fraction of the rays which passed through a circular aperture S in a metal screen was measured by passing them into an ionization chamber Q containing air or some other gas, and finding the conductivity produced. The ratio of the conductivity with the scattering plate at P to the conductivity without any scattering plate is approximately equal to  $n/n_0$ . The scattering angle for the rays passing through the aperture S is less than  $\tan^{-1}(a/d)$ , where  $a$  is the radius of the aperture and  $d$  its distance from the scattering plate.

The results obtained by Crowther and other physicists agree approximately with the equation  $n/n_0 = 1 - \varepsilon - \phi^2/kt$ , where  $k$  is a constant. With different sized apertures the thickness of the plate required to produce a given value of  $n/n_0$  is proportional to  $\phi^2$ , in agreement with the theory. The following table gives some results obtained by Crowther and Schönland with  $\phi = 0.11$  for the scattering of  $\beta$ -rays by gold.

$t$ .	$n/n_0$ (Observed).	$t \log \frac{n_0}{n_0 - n}$	$n/n_0$ (Calculated).
1	0.86	0.85	0.89
2	0.67	0.96	0.67
3	0.52	0.96	0.52
4	0.46	1.07	0.43
5	0.38	1.03	0.36

The first column gives the number of gold foils used as the scattering plate, which is proportional to  $t$ . If  $\frac{n}{n_0} = 1 - \epsilon^{-\phi^2/kt}$ , then

$$\frac{\phi^2}{k} = t \log \frac{n_0}{n_0 - n},$$

so that the numbers in the third column should be constant according to the theory. The last column gives the calculated values of  $n/n_0$ , taking  $\phi^2/k$  equal to 0.97, which is the mean of the numbers in the third column. Crowther and Schönland's results give  $k = 680$  for gold. Now  $k = \pi \mathcal{N} R^2 \phi_1^2$ , so that we can calculate  $\phi_1^2$  if we know  $\mathcal{N}$  and  $R$ . The number of atoms of gold per cubic centimetre is  $6 \times 10^{22}$ , and we may assume  $R = 10^{-8}$  cm., so that we get

$$\phi_1^2 = \frac{680}{\pi \times 6 \times 10^{22} \times 10^{-16}} = 3.6 \times 10^{-5}.$$

According to this the root mean square deviation of the  $\beta$ -rays by an atom of gold was only  $6 \times 10^{-3}$  or about  $\frac{1}{3}$  of a degree. The  $\beta$ -rays for which the above results were obtained had a velocity corresponding to a fall of potential of  $4.6 \times 10^5$  volts.

If we assume that the atoms consist of a positively charged nucleus surrounded by a number of electrons it is possible to calculate the average deviation of  $\beta$ -rays passing through such atoms, and the results obtained are consistent with the observed average deviations. For the details of such calculations the original papers may be consulted.

### 11. Absorption of $\beta$ -rays.

When  $\beta$ -rays from a layer of radioactive material are absorbed by placing a thin sheet of metal over the layer, so that the rays fall on the sheet in all directions, then the intensity of the rays getting through is proportional approximately to  $\epsilon^{-\mu t}$ , where  $\mu$  is a constant and  $t$  the thickness of the sheet. The intensity may be measured by the ionization produced by the rays. The constant  $\mu$  is roughly speaking proportional to the density  $\rho$  of the material of the sheet, as with cathode rays. The following are some values of  $\mu/\rho$  for the  $\beta$ -rays from uranium.

Substance.	$\mu/\rho$
Silver .. ..	7.3
Aluminium .. ..	4.1
Lead .. ..	9.75
Sulphur .. ..	4.5
NaCl .. ..	4.7
KI .. ..	7.8

For the  $\beta$ -rays from radium-E  $\mu/\rho$  varies from 15.8 for carbon to 22.1 for tin. The following table gives, in centimetres, the thickness of aluminium penetrated by  $\beta$ -rays which describe a circle of radius  $R$  in a magnetic field of strength  $H$  perpendicular to the plane of the circle.

$HR$	=	1,380	2,535	3,790	5,026	7,490	11,370
Thickness	=	0.018	0.124	0.279	0.440	0.785	1.36

When  $\beta$ -rays pass through thin sheets of matter their velocity is decreased. The change in the product  $RH$  due to a sheet weighing 0.01 gm. per square centimetre is given approximately by

$$\Delta(RH) = K \frac{c^3}{v^3},$$

where  $v$  is the velocity of the rays,  $c$  that of light,  $\Delta(RH)$  the decrease in  $RH$  and  $K$  a constant. The constant  $K$  is about 35 for mica, from 23 to 32 for tin, and about 28 for gold.

#### $\alpha$ -RAYS

### 12. Ratio of Charge to Mass.

The properties of  $\alpha$ -rays will now be considered.  $\alpha$ -rays are rays emitted by many radioactive bodies; they are much less penetrating than the  $\beta$ -rays. They are deflected by electric and magnetic fields in the opposite direction to  $\beta$  and cathode rays, and are found to carry a positive charge.

The ratio of the charge  $e$  to the mass  $m$  of  $\alpha$ -rays has been found by a method similar to that used by Bucherer for  $\beta$ -rays. The  $\alpha$ -rays were passed between two parallel plates very near together and a potential difference was maintained between the plates. The apparatus was put in a uniform magnetic field perpendicular to the electric field between the plates and to the path of the rays. By properly adjusting the field strengths the force on the rays due to the electric field could be made equal and opposite to that due to the magnetic field, so that the rays were not deflected and could pass between the plates. In this case we have  $Fe = Hev$ , where  $F$  is the electric and  $H$  the magnetic field strength, so that the velocity  $v$  of the rays is given by  $v = F/H$ . After passing between the plates the rays emerge into the uniform magnetic field and describe circular paths in it of radius  $r$  given by  $mv^2/r = Hev$ . The rays fell on a photographic plate on which their deflection could be measured, and the radius  $r$  could be calculated from the deflection and the dimensions of the apparatus. In this way it was found that  $e/m = 4823$  e.m.u. per gramme, and that  $v$  is between  $1.4 \times 10^9$  and  $2.2 \times 10^9$  cm. per second for the  $\alpha$ -rays from different radioactive bodies.

### 13. Counting of $\alpha$ -rays. Scintillations. Ionization.

A remarkable property of  $\alpha$ -rays was discovered by Sir William Crookes. He found that when  $\alpha$ -rays fall on a screen coated with powdered zinc-blende the screen emits minute flashes of light easily visible in a low-power microscope. These flashes or scintillations are each due to the impact of one  $\alpha$ -ray particle, so that when the rays are sufficiently feeble it is possible by counting the scintillations to determine the number of  $\alpha$ -ray particles falling on the screen. Other fluorescent substances besides zinc-blende may be used—diamond, for example, makes a very good screen.

$\alpha$ -rays ionize gases strongly, and the ionization due to a single  $\alpha$ -ray particle can be easily detected, which gives another method of counting the particles. The apparatus used by Rutherford to count the  $\alpha$ -rays emitted by radium-C is shown in fig. 11. The radium-C was deposited on a small cone at S in a long glass tube EF provided with a stop-cock at T. At D there was a metal diaphragm having a small circular hole in it through which some of the  $\alpha$ -rays from S

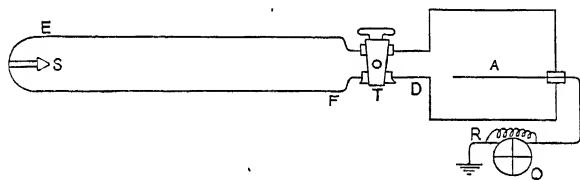


Fig. 11

passed into a metal box in which there was an insulated electrode A. The potential difference between the box and the electrode, and the gas pressure in the apparatus, were adjusted so that an electron set free in the box produced a large number of ions by collisions with the gas molecules. In this way the ionization due to a single  $\alpha$ -ray entering the box through D was made sufficient to cause an easily observable deflection of a quadrant electrometer connected to the insulated electrode A. The electrode A was also connected to the earth through a very high resistance R, so that the deflection due to an  $\alpha$ -ray was not permanent but rapidly disappeared. The number of deflections due to  $\alpha$ -rays was counted, and so the number passing through the hole at D in a known time found. The total number emitted by the radium-C could then be calculated by assuming the rays to be emitted equally in all directions.

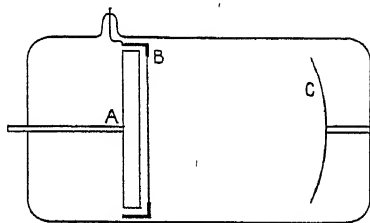


Fig. 12

#### 14. Charge carried by $\alpha$ -rays.

The charge carried by the  $\alpha$ -rays emitted by radium-C was found with the apparatus shown in fig. 12. The radium-C was deposited on a plate C supported in a highly exhausted glass tube. The  $\alpha$ -rays from C were received on an insulated electrode A after passing through a sheet of thin foil B and another sheet of thin foil covering the front of A. When  $\alpha$ -rays strike a metal surface they cause the emission of electrons which leaves a positive charge on the metal in addition to the charge carried by the  $\alpha$ -rays. To prevent these electrons from escaping from the electrode A it was covered with the thin foils which

allowed the  $\alpha$ -rays to pass through but stopped the electrons. The apparatus was also put in a strong magnetic field perpendicular to the plane of the paper which caused the electrons to describe small circular or spiral paths and so helped to prevent them from escaping. The charge received by the electrode A in a known time was measured with a quadrant electrometer. The total charge on the  $\alpha$ -rays emitted by the radium-C in a known time could then be calculated by assuming the rays to be emitted equally in all directions from every point on the plate at C. The amount of radium-C on the plate C was compared with that used in the previous experiment by comparing the intensities of the  $\gamma$ -rays emitted by the two deposits. In this way it was found that an amount of radium-C equal to that contained in 1 gm. of radium in radioactive equilibrium with its products emits  $3.4 \times 10^{10}$   $\alpha$ -rays per second and that the charge on these rays is 31.6 electrostatic units. The charge carried by one  $\alpha$ -ray particle is therefore  $9.3 \times 10^{-10}$  electrostatic units. Since  $e/m$  is 4823 electromagnetic units per gramme it follows that the mass of one  $\alpha$ -ray particle is given by

$$m = \frac{9.3 \times 10^{-10}}{3 \times 10^{10} \times 4823} = 6.42 \times 10^{-24} \text{ gm.}$$

The mass of one hydrogen atom is  $1.663 \times 10^{-24}$  gm., so it appears that the mass of one  $\alpha$ -ray particle is nearly four times that of one hydrogen atom.

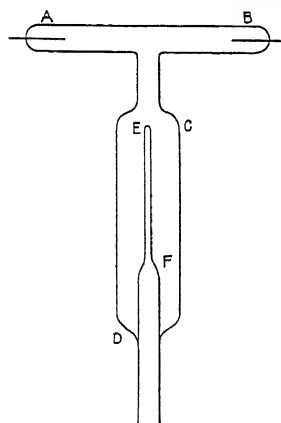


Fig. 13

### 15. $\alpha$ -rays are Charged Helium Atoms.

The ratio  $e/m$  for positively charged hydrogen atoms or hydrogen ions in solution is 9650, which is just double that found for  $\alpha$ -rays. Also the charge  $9.3 \times 10^{-10}$  on one  $\alpha$ -ray is nearly double the charge on one gaseous ion or electron. We conclude that the charge on one  $\alpha$ -ray is  $2e$  and that the mass of one  $\alpha$ -ray particle is four times that of one hydrogen atom. Now the mass of one helium atom is known to be four times that of one hydrogen atom, which suggests that  $\alpha$ -rays may be charged helium atoms.

That the  $\alpha$ -rays are charged helium atoms was shown very clearly by an experiment due to Rutherford. The apparatus used is shown in fig. 13. A small discharge tube AB was connected to a tube CD and highly exhausted. The tube CD was sealed on to a tube EF, the end of which was drawn out into a narrow very thin walled tube. Some radium emanation was pumped into the narrow part of EF, and the  $\alpha$ -rays emitted by the emanation passed through the thin glass

walls into CD and AB. A discharge could then be passed through AB and the light emitted gave the spectrum of pure helium.

The helium atom is supposed to consist of a positively charged nucleus having a charge  $2e$  and two electrons each having a charge  $-e$ . The  $\alpha$ -ray particles are therefore supposed to be simply nuclei of helium atoms.

### 16. Range and Velocity of $\alpha$ -rays.

When  $\alpha$ -rays from a very thin film of some radioactive body are passed through a gas it is found that they all travel very nearly the same distance before they are stopped. This distance is called the range of the  $\alpha$ -rays. The range in a gas is found to be inversely as the gas pressure.

The range may be determined by means of the apparatus shown in fig. 14. A spherical glass bulb AA' is coated on the inside with a thin film of finely powdered zinc-blende, and the radioactive material which emits the  $\alpha$ -rays is deposited on a small sphere B which is supported at the centre of the bulb on a rod attached to a stopper S. The gas pressure in the bulb can be varied by means of a side tube T which leads to a pump and manometer. The gas pressure is adjusted until the  $\alpha$ -rays from B just reach the bulb, as shown by the scintillations produced on the zinc-blende film. The range is then equal to the radius of the bulb, and the range in the gas at 760 mm. pressure can be calculated since the range is inversely as the pressure. The range of the  $\alpha$ -rays from radium-C is 7.0 cm. in air at 760 mm., and that of the rays from radium is 3.5 cm. The velocity of  $\alpha$ -rays diminishes as they pass through a gas. Rutherford found the velocity of  $\alpha$ -rays which had passed through different thicknesses of air by measuring the magnetic deflection of the rays, and found that

$$v^3 = A(R - x),$$

where  $v$  is the velocity of the rays,  $A$  a constant,  $R$  the range of the rays, and  $x$  the distance travelled from the source. Thus, when  $x = R$ ,  $v = 0$  as we should expect. Differentiating the above equation with respect to the time  $t$  we get

$$\frac{dv^2}{dt} = -\frac{2}{3}A,$$

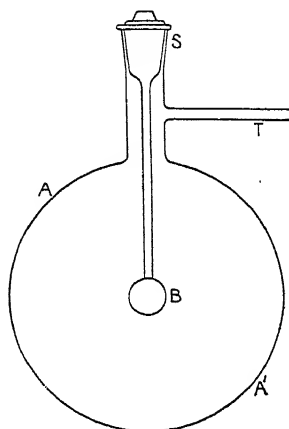


Fig. 14

so that it appears that the rate of loss of kinetic energy as the rays move through a gas is constant. Also  $\frac{dv^2}{dx} = -\frac{2}{3}\frac{A}{v}$ , so that the energy lost per unit distance moved is inversely as the velocity. We should expect the ionization produced to be proportional to the loss of energy, and therefore the ionization per centimetre to be inversely as  $(R - x)^{1/3}$ .

### 17. Ionization by $\alpha$ -rays. Stopping Power.

The ionization produced by  $\alpha$ -rays along the range was investigated by Bragg with the apparatus shown in fig. 15. The  $\alpha$ -rays were emitted by a film of radioactive material on a horizontal disc R, which carried a grating made of vertical metal plates so that only those  $\alpha$ -rays moving in nearly vertical directions could get out. In this way a nearly parallel vertical beam of the rays was obtained. The rays passed through the gas in a box AB and through a horizontal plate of fine wire gauze at C. Just above the gauze there was an insulated electrode D. The rays ionized the gas between C and D, and this ionization was measured by connecting D to a quadrant electrometer. Sufficient potential difference was maintained between C and D to saturate the current. By moving the source R up and down, the ionizations due to the rays at different distances from the source could be compared.

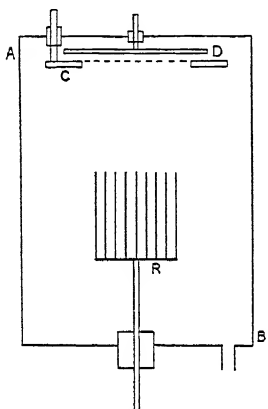


Fig. 15

It was found that the ionization increased with the distance from the source to a maximum when the distance was nearly equal to the range of the rays and then rapidly fell to zero. The ionization was approximately proportional to  $(R - x)^{-1/3}$  and therefore to the kinetic energy lost by the rays. The range in any gas is inversely as the pressure, so that the mass of gas per unit area in a layer of thickness equal to the range is independent of the gas pressure. This mass is  $R\rho$  where  $\rho$  is the density of the gas. The *stopping power* of the gas is defined as  $R\rho$  for air divided by  $R\rho$  for the gas.

If  $\alpha$ -rays fall normally on a very thin sheet of any solid and pass through, their velocity and range in air or other gas are diminished. The decrease of the range may be found with the apparatus shown in fig. 15. If the thin sheet is put over the source R so that the rays pass through it, the decrease of the distance at which the maximum ionization is obtained is equal to the decrease of the range. The decrease of range due to a thin sheet is nearly proportional to the mass  $m$  per

unit area of the sheet. If the weight of an atom of the material of the sheet is  $A$ , then  $m/A$  is the number of atoms per unit area in the sheet. Let  $\Delta R$  denote the decrease of range in air of density  $\rho$ , and  $A'$  the mean weight of the air atoms. Then  $\rho \Delta R/A'$  is the number of air atoms per unit area in a layer which diminishes the range by the same amount as the sheet. The *atomic stopping power* of the atoms in the sheet is defined as the ratio of the number of air atoms  $\rho \Delta R/A'$  to the number  $m/A$  of atoms per unit area in the sheet. The stopping power is therefore equal to  $\rho A \cdot \Delta R/mA'$ . The atomic stopping power of different elements is found to be roughly proportional to the square root of the atomic weight. For example, the atomic stopping power of silver is 3.1, and the square root of the ratio of the atomic weights of silver and air is  $\sqrt{108/14.4} = 2.74$ , which is not far from 3.1.

When  $\alpha$ -rays go through thin metal sheets most of them go nearly straight through, but a small fraction is deviated through large angles. This scattering of  $\alpha$ -rays has been investigated by Rutherford, Geiger, Chadwick, and others with very important results.

### 18. Single Scattering of $\alpha$ -rays.

The theory of the scattering of  $\beta$ -rays discussed earlier in this chapter was based on the assumption that the deviations were the resultants of a large number of small deviations, the average deviation due to passing through one atom being only a rather small fraction of a degree. Such scattering is called *multiple scattering*. When  $\alpha$ -rays are passed through matter the great majority are deviated through small angles, and these small deviations obey the same laws as the deviations of  $\beta$ -rays, i.e. the number  $n$  out of a total number  $n_0$  deviated through angles less than  $\phi$  is given by  $n/n_0 = 1 - e^{-\phi^2/\bar{\phi}^2}$ , where  $\bar{\phi}^2$  is the average value of the squares of the deviations. According to this formula the number deviated through angles much greater than the average deviation is quite negligible. Thus if  $\phi/\bar{\phi}$  is equal to 3 then  $n/n_0 = 1 - 0.00012$ . It is found, however, that for large values of  $\phi$  the number of  $\alpha$ -rays scattered does not fall off with  $\phi$  as rapidly as the theory of multiple scattering indicates, so that there is an appreciable number even with very large values of  $\phi$ . To explain this Rutherford proposed the theory of *single scattering*, according to which the large deviations are due to single collisions with atoms and not to the summation of a large number of small deviations. Suppose  $\alpha$ -rays to pass through a sheet consisting of a single layer of atoms, so that each ray passes through one atom, and suppose further that all but a very small fraction, say 1/1000, are only deviated through very small angles. Now let the rays pass through a second similar sheet. The chance that a particular  $\alpha$ -ray will be deviated through more than a small angle by both sheets is only 1 in 1,000,000 and so is negligible. Even for a sheet consisting of 100 layers of atoms the chance that a ray is deviated once through more than a small angle is only about 1 in 10, and therefore the chance that a ray suffers two large deviations is about 1 in 100. In this way it is easy to see that the single-scattering theory should apply to the rays scattered through large angles by thin sheets, provided the total number scattered through large angles is only a small fraction of the whole number. According to this the scattering of  $\alpha$ -rays through large angles by thin sheets will be the same as if the rays had all passed through one atom, and so will depend on the field of force inside the atom. To explain the large deviations an intense field in the atom is required,

and this led Rutherford to propose his nucleus theory of the atom. According to this theory an atom consists of a positively charged nucleus surrounded by a number of electrons. In a neutral atom the positive charge on the nucleus is equal to the negative charge on the electrons, so that the charge on the nucleus is always an exact multiple of the charge  $e$ , which is a positive charge equal to the negative charge on one electron. Owing to the small mass of electrons it is clear that the  $\alpha$ -rays cannot be deviated appreciably by them, so that the large deviations must be produced when an  $\alpha$ -ray passes very close to a nucleus.

In fig. 16 let ABC be the path of an  $\alpha$ -ray passing a nucleus at N. Let AD

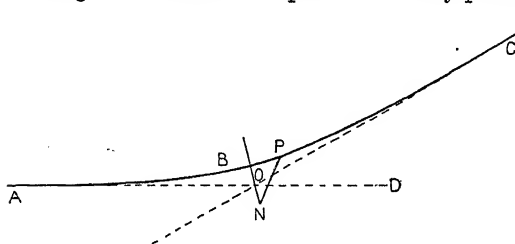


Fig. 16

be the path the ray would have followed in the absence of the nucleus, and let  $\angle COD$  be the deviation  $\phi$ . Let  $p$  be the length of the perpendicular on AD from N. We shall suppose the nucleus fixed in position at N. Let the co-ordinates of a point P on the path be  $NP = r$  and  $\angle PNO = \theta$ . The angular

momentum of the  $\alpha$ -ray about N is  $mv_0 p = h$ , a constant, where  $v_0$  is the initial velocity of the ray along AD, and  $m$  is the mass of the ray.

The kinetic energy of the ray is given by

$$\frac{1}{2}mv_0^2 - \frac{1}{2}mv^2 = 2Ee/r,$$

where  $v$  is the velocity of the ray when at a distance  $r$  from the nucleus,  $2e$  is the charge of the  $\alpha$ -ray, and  $E$  is the charge of the nucleus. The electrical potential due to the nucleus is  $E/r$ , so that the ray loses kinetic energy equal to  $2Ee/r$  as it approaches the nucleus.

Now  $v^2 = \dot{r}^2 + (r\dot{\theta})^2$ , where  $\dot{r} = dr/dt$  as usual. But  $\dot{r} = \frac{dr}{d\theta}\dot{\theta}$ ; so that

$$v^2 = \left\{ \left( \frac{dr}{d\theta} \right)^2 + r^2 \right\} \dot{\theta}^2.$$

We have also  $mv_0 p = h = mr^2\dot{\theta}$ , so that we get

$$v^2 = \left\{ \left( \frac{dr}{d\theta} \right)^2 + r^2 \right\} \frac{h^2}{m^2 r^4}.$$

Now let  $r = u^{-1}$ , so that  $\frac{dr}{d\theta} = -u^{-2} \frac{du}{d\theta}$ , and

$$v^2 = \left\{ u^{-4} \left( \frac{du}{d\theta} \right)^2 + u^{-2} \right\} \frac{h^2}{m^2} u^4.$$

Hence

$$\left( \frac{du}{d\theta} \right)^2 + u^2 = \frac{m^2}{h^2} \left\{ -\frac{4Ee}{m} u + v_0^2 \right\}.$$

The solution of this equation (cf. p. 329) is

$$u = A(\varepsilon \cos \theta - 1),$$

where  $A$  and  $\varepsilon$  are constants. Substituting this in the differential equation we find

$$A = \frac{Ee}{Tp^2} \text{ and } \varepsilon^2 - 1 = \left( \frac{Tp}{Ee} \right)^2,$$

where  $T = \frac{1}{2}mv_0^2$  is the initial kinetic energy of the  $\alpha$ -ray.

In fig. 16, OA and OC are asymptotes to the path. Let  $\angle AOB = \angle BOC = \psi$ ,  $\angle COD = \varphi$ .

When  $r$  is very large we have  $u = 0$ , so that  $\varepsilon \cos \psi = 1$ , and we have then  $\varphi + 2\psi = \pi$ , or  $\frac{1}{2}\varphi = \frac{1}{2}\pi - \psi$ , and therefore  $\sin \frac{1}{2}\varphi = 1/\varepsilon$ , and  $\cot \frac{1}{2}\varphi = \sqrt{\varepsilon^2 - 1}$ , or

$$\cot\left(\frac{\varphi}{2}\right) = \frac{Tp}{Ee}.$$

Suppose now that a large number  $n$  of  $\alpha$ -rays pass through a thin plate of thickness  $t$  and let the number of nuclei in the plate be  $N$  per unit volume. The total path of the rays in the plate is  $nt$ , so that the number of cases in which the undeviated path of a ray comes nearer to a nucleus than a distance  $p$  will be equal to the number of nuclei in a cylinder of length  $nt$  and cross-section  $\pi p^2$ . The number of such cases is therefore  $\pi ntp^2N$ , and the number of rays which are deviated through an angle greater than  $\varphi$  is

$$\pi n t N \left(\frac{Ee}{T}\right)^2 \cot^2\left(\frac{\varphi}{2}\right).$$

The fraction of the rays deviated through angles greater than  $\varphi$  is therefore

$$\pi N t \left(\frac{Ee}{T}\right)^2 \cot^2\left(\frac{\varphi}{2}\right).$$

We have not taken into account the charges on the electrons, which, of course, must limit the electric field due to the nucleus. However, the electrons are believed to be at distances from the nucleus of the order of  $10^{-8}$  cm., and the values of  $p$  required to give large values of  $\varphi$  are much smaller than this, so that for the large deflections the field near the nucleus may be taken to be equal to  $E/r^2$ , as if the electrons were not present.

The number of  $\alpha$ -rays scattered through different angles by thin plates of gold and other metals was found by passing a narrow nearly parallel beam of the rays through the plate and counting the scintillations produced on a small screen by the scattered rays. The small screen could be set up so as to receive the rays scattered through any desired angle. The total number of rays falling on the plate was also determined.

The results obtained agreed very closely with the single-scattering theory. The fraction of the rays scattered through an angle greater than  $\varphi$  was found to be proportional to the thickness of the scattering plate, inversely proportional to the square of the energy  $T$  of the rays, and proportional to  $\cot^2(\varphi/2)$ . From the values found for the fraction scattered it was possible to calculate the nuclear charge  $E$  for the atoms of the plate. It was found in this way, for example, that the nuclear charge for gold atoms is equal to  $(79 \pm 0.5)e$ , in good agreement with the theory that the nuclear charge is equal to the product of the electronic charge by the atomic number, for the atomic number of gold is 79. Similar results were obtained with several other metals. These experiments therefore afford very strong support for Rutherford's nucleus theory of the atom.

## 19. $\alpha$ -rays and Fast Hydrogen Atoms.

When an  $\alpha$ -ray collides with a gas molecule we should expect it to be possible for the molecule to acquire a large velocity comparable with that of the  $\alpha$ -ray. It is found that when  $\alpha$ -rays are passed through hydrogen gas some hydrogen atoms acquire large velocities and are able to produce scintillations like  $\alpha$ -rays.  $\alpha$ -rays which have a range

of 24 cm. in hydrogen at 760 mm. produce some rapidly moving positively charged hydrogen atoms which have a range of 58 cm. in the hydrogen. Similar long-range particles, apparently positively charged hydrogen atoms, have also been obtained by Rutherford by passing  $\alpha$ -rays through thin plates containing certain elements of small atomic weight but no hydrogen. It is supposed that the  $\alpha$ -rays collide with the atomic nuclei and knock a positively charged hydrogen atom out of them. This has been done with boron, nitrogen, fluorine, sodium, aluminium, and phosphorus. This phenomenon is discussed in the chapter on Constitution of Atoms.

The method due to C. T. R. Wilson of making visible the tracks of  $\alpha$ - and  $\beta$ -rays through gases and the results obtained are described in the chapter on Gaseous Ions.

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## CHAPTER X

# Positive Rays

### 1. Nature of Positive Rays.

Positive rays are positively charged atoms or molecules which have acquired a high velocity in an electric field. They are produced in the electric discharge in gases at low pressures and in other ways. If a discharge is passed between two electrodes in a gas at a low pressure the gas is ionized, that is, electrons are removed from the gas molecules, which therefore become positively charged. These positive ions move towards the negative electrode, and when the potential difference is large, say 50,000 volts, they acquire a high velocity. If a small hole is bored through the negative electrode, some of the positive ions pass through the hole and form a beam of positive rays. They can be deflected by transverse magnetic and electric fields. The deflections are much smaller than those of cathode rays or electrons and are in the opposite direction, showing that the rays are positively charged.

These rays have been investigated by Wien, J. J. Thomson, Aston, and others, and very important results have been obtained. The positive rays cause gases through which they pass to emit light, and they produce phosphorescence when they fall on solid bodies. The mineral willemite, a silicate of zinc, phosphoresces brightly when exposed to positive rays and was found by J. J. Thomson to be one of the best substances for locating a beam of the rays. They affect a photographic plate like light, and so measurements of their deflections are usually made by photographing them.

### 2. Experiments of J. J. Thomson.

A discharge tube used by J. J. Thomson is shown in fig. 1. It consists of a large glass bulb BB of about 1500 c. c. capacity, having a small electrode sealed in at D. The cathode C consists of an iron tube about 7 cm. long with an aluminium cap on it at C. Along the axis of this tube a very fine straight copper tube is mounted, through which the rays pass into a conical tube K where they fall on a willemite screen or photographic plate P. The narrow beam of rays is passed between two insulated iron blocks E and F which can be magnetized by the poles G, H of a large electromagnet, so producing a magnetic

between E and F perpendicular to the beam of rays. The blocks E and F can be connected to a battery so as to produce an electric field between them in the same direction as the magnetic field. The blocks B and tube K are connected to pumps by means of which they can be exhausted.

A nearly perfect vacuum is maintained in K, and the gas to be examined is slowly admitted to B through a narrow tube, the pressure in B being kept constant at a suitable small value by the pumps. The electrode D is charged positively by a large induction coil and the cathode C connected to the earth.

When the pressure in B is low the potential difference required to produce a discharge in the bulb is large and positive rays are produced, some of which pass through the narrow tube and give a small spot on the plate P. When the magnetic and electric fields between E and F are produced the rays are deflected and distributed on the plate P along parabolic curves, the positions of which depend on the charges, masses, and velocities of the rays.

### Theory of the Positive Ray Parabolas.

The force on a particle of mass  $m$ , carrying an electric charge  $e$ , moving with velocity  $v$  in a magnetic field  $H$  perpendicular to  $v$ , is  $Hev$  and is perpendicular to  $v$ . If then the particle travels a distance  $d$  in the magnetic field it will acquire a velocity  $u$  perpendicular to the field given by  $mu = Hevd/v$ , so that  $Hed/m$ . The deflection of the particle after travelling a distance  $l$  beyond the field will be approximately  $y = lu/v = Hedl/mv$ .

The force on the particle in an electric field  $F$  is  $Fe$ , so that while travelling a distance  $d$  in an electric field  $F$  perpendicular to  $v$  it will acquire a velocity in the direction of  $F$  given by  $mw = Fed/v$ . The deflection  $z$  after going a further distance  $l$ , where  $F = 0$ , is given by  $z = lw/v = Fedl/mv^2$ .

J. J. Thomson's apparatus  $H$  and  $F$  were both perpendicular to the faces of the blocks  $E$  and  $F$ , and both extended over the same distance  $d$ , so that the electric and magnetic deflections were at right angles to each other and equal to  $z$ . The positive rays having a given value of  $e/m$  but different velocities therefore fall on a parabola on the plate P, the equation of which is got by eliminating  $v$  from

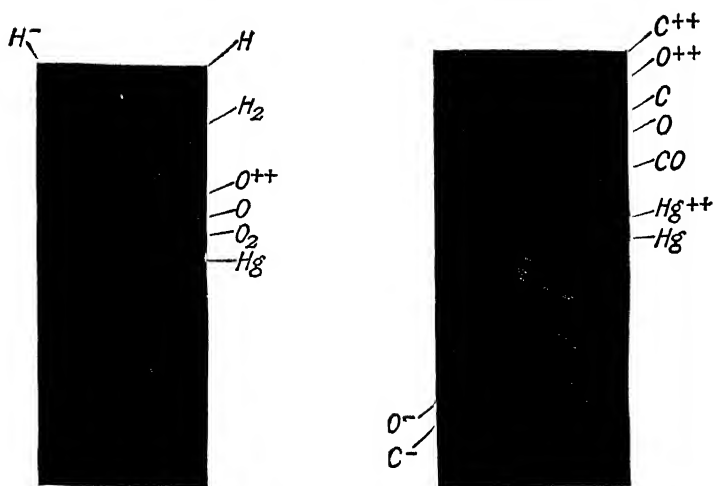
$$y = \frac{Hedl}{mv} \text{ and } z = \frac{Fedl}{mv^2}.$$

The equation of the parabola is therefore

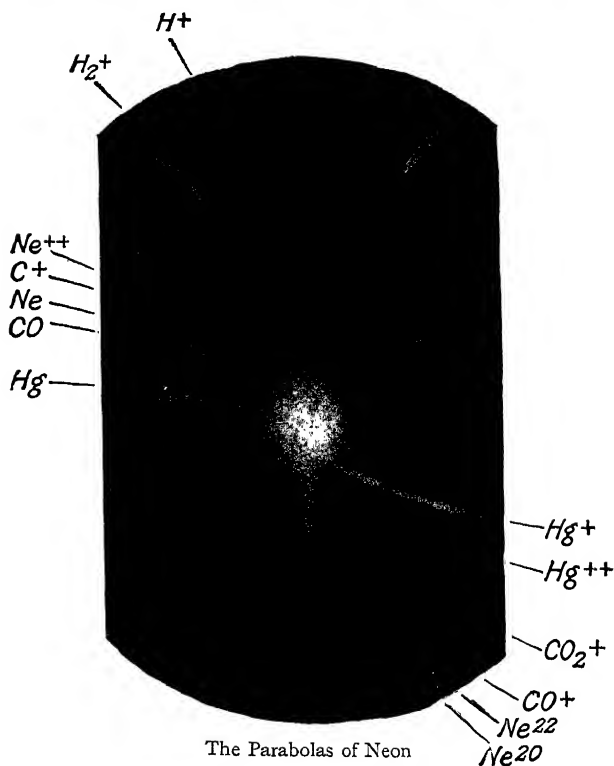
$$\frac{Fedl}{mz} = \left( \frac{Hedl}{ym} \right)^2,$$

$$y^2 = \frac{H^2}{F} \frac{edl}{m} z.$$





Photographs of Typical Positive Ray Parabolas



The Parabolas of Neon

Fig. 2

After Aston, *Isotopes* (Arnold).

By measuring  $y$  and  $z$  for points on the parabola,  $e/m$  can be determined when  $H$ ,  $F$ ,  $d$ , and  $l$  are known.

When a mixture of gases is put in the discharge tube and the positive rays from it examined in this way a number of distinct parabolas is obtained on the plate P, one for each value of  $e/m$  of the positive rays produced.

The ratio of  $y$  to  $z$  is given by

$$\frac{y}{z} = \frac{H}{F} v,$$

so that all rays having the same velocity but different values of  $e/m$  fall on a straight line passing through the origin ( $y = 0, z = 0$ ) and inclined to the  $z$  axis at an angle  $\theta$  given by

$$\tan \theta = \frac{H}{F} v.$$

Fig. 2 shows several sets of parabolas obtained on photographic plates by J. J. Thomson. The letters at the ends of the parabolas indicate the atom or molecule to which the parabola is attributed. If the charge on the atom is equal to that on an atom which has lost one electron or  $+e$ , this is either not indicated or indicated by one plus sign, and if the charge is  $2e$  this is indicated by two plus signs; thus  $\text{Hg}^{++}$  indicates the parabola attributed to mercury atoms each having a charge  $2e$ .

If  $m_H$  is the mass of one hydrogen atom,  $m_R$  the mass of one positive ray, and  $e_H$  and  $ne_H$  are the charges they carry, then  $e_H/m_H = 9650$  electromagnetic units per gramme, so that if  $ne_H/m_R$ , which is  $e/m$  for the positive rays, is equal to  $x$ , then

$$\frac{9650}{x} = \frac{m_R}{nm_H}.$$

If the positive ray consists of  $N$  atoms of atomic weight  $A$ , then  $NA = m_R/m_H$  very nearly, so that  $x = 9650n/NA$ .

For example, in the case of mercury,  $A = 200$ , and we get the following possible values of  $e/m$  or  $x$  for single mercury atoms:

$n$ .		$e/m = x$ .
1	....	48.25
2	....	96.5
3	....	144.75
4	....	193.0
5	....	241.25
6	....	289.50
7	....	337.75

It was found that with mercury vapour seven distinct parabolas could be obtained corresponding to the values of  $e/m$  given above. Thus it appears that mercury atoms can lose from one to seven electrons in the discharge tube.

In hydrogen, parabolas corresponding to  $n = 1, N = 1$ ;  $n = 1, N = 2$ ; and  $n = 1, N = 3$  were obtained.

The parabolas of many elements and compounds have been observed and the corresponding values of  $e/m$  are always such as can be readily explained by supposing the rays to consist of an atom or molecule of the substance in question which has lost one or more electrons.

#### 4. Have All the Atoms of a Given Element the Same Mass?

This method of positive ray analysis gives a definite answer to the question whether all the atoms of a given element have equal

masses or not. If well defined parabolas are obtained they must be due to particles all having the same value of  $e/m$  and so presumably of  $m$ . If the masses of the atoms of an element varied between certain limits above and below a mean value equal to that corresponding to the chemical atomic weight, then well defined parabolas would not be obtained but more or less broad diffuse ones. It is clear, therefore, that this is not the case. However, the possibility remains that the atoms of a given element are not all of the same mass, but that there are two or more possible masses each of which gives a well defined parabola. It is found that such is the case. J. J. Thomson obtained two parabolas with neon corresponding to atomic weights 20 and 22, whereas the accepted atomic weight of neon is 20.2. Aston and G. P. Thomson found that lithium also gives two parabolas corresponding to atomic weights 6 and 7, whereas the chemical atomic weight is 6.94. G. P. Thomson has examined several light elements by the parabola method. He found beryllium to be a simple element.

### 5. Aston's Method of Positive Ray Analysis.

A new method of positive ray analysis was invented by Aston which enables  $e/m$  to be determined with much greater accuracy than by the parabola method. Aston's apparatus is shown in fig. 3.

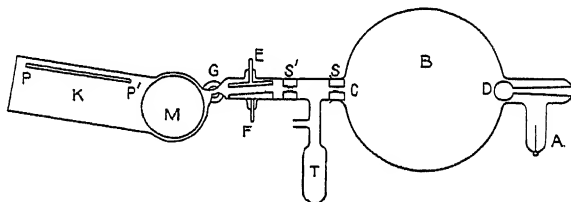


Fig. 3

The discharge producing the positive rays takes place in a large bulb B between a concave cathode C and an anode A. The cathode rays from C fall on a quartz bulb D and are thus prevented from fusing the glass wall of the discharge tube. In the cathode C there is a narrow slit S through which the positive rays pass. A second slit at S' allows a narrow beam of the rays from S to pass. The space between S and S' is kept highly exhausted by means of a bulb T containing charcoal cooled in liquid air. The beam of rays passes between two parallel metal plates E and F between which an electric field is maintained. The rays are deflected by this field and the narrow beam is spread out into a broad band. Part of this band is allowed to pass through an adjustable slit G. The rays then pass between the poles of a magnet M and are deflected by the magnetic field in the opposite direction to the previous electric deflection. Finally, they pass into

a box K and fall on a photographic plate PP'. This arrangement causes all rays having the same value of  $e/m$  to strike the plate at the same point, so that lines are obtained on the plate one for each value of  $e/m$ . The series of lines on the plate is called a "mass spectrum" and the apparatus a "mass spectrograph". The box K and the rest of the apparatus except the bulb B must be highly exhausted, since any appreciable amount of gas stops the positive rays. The gas to be examined is slowly admitted to B and the pressure kept constant by pumps as in J. J. Thomson's apparatus.

## 6. Theory of Aston's Mass Spectrograph.

Let the angle through which the rays are deflected by the electric field be  $\theta$  and by the magnetic field  $\phi$ . Also let  $b$  be the distance from the middle of the electric field to the middle of the magnetic field.

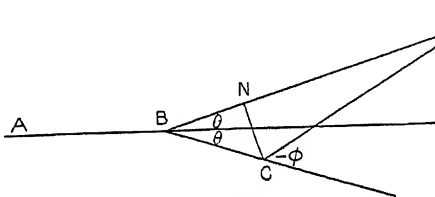


Fig. 4

Consider two rays for which  $e/m$  is the same but which have velocities  $v$  and  $v + dv$  respectively. Then we have approximately

$$\theta = \frac{Fed}{mv^2}, \text{ and } \phi = \frac{Hed'}{mv},$$

as in J. J. Thomson's apparatus, where  $d$  is the length of the electric and  $d'$  that of the magnetic field. Hence

$$d\theta = -2 \frac{Fed}{m} \frac{dv}{v^3}, \quad d\phi = -\frac{Hed'}{mv^2} dv,$$

so that 
$$\frac{d\theta}{\theta} = -2 \frac{dv}{v}, \text{ and } \frac{d\phi}{\phi} = -\frac{dv}{v},$$

and therefore 
$$d\phi = \frac{\phi}{2\theta} d\theta.$$

The separation of the two rays at the magnetic field is  $bd\theta$ , and at a distance  $r$  farther on it is  $bd\theta + r(d\theta + d\phi)$  or  $d\theta \left\{ b + r \left( 1 + \frac{\phi}{2\theta} \right) \right\}$ .

This is zero when 
$$b + r \left( 1 + \frac{\phi}{2\theta} \right) = 0,$$

or when 
$$r = \frac{-b}{1 + \phi/2\theta}.$$

This requires  $\phi$  to be negative and greater than  $2\theta$  for  $r$  to be positive.

In fig. 4 let AB be the original direction of the positive rays and let them be electrically deflected at B through the angle  $\theta$  along BC.

Let the magnetic deflection take place at C through the angle  $-\phi$  greater than  $2\theta$  so that they go along CD. Draw BND making an angle  $\theta$  with AB produced and let CD and BND meet at D. Then the angle at D is equal to  $-(\phi + 2\theta)$ , so that

$$NC = b \sin 2\theta = -r \sin(\phi + 2\theta),$$

where  $b = BC$ , and  $r = CD$ . If  $\theta$  and  $\phi$  are small angles then approximately

$$2b\theta = -r(\phi + 2\theta),$$

or

$$r = \frac{-b}{1 + \phi/2\theta}.$$

Thus we see that rays having the same value of  $e/m$  but slightly different velocities will all cross BND at the same point. In Aston's apparatus the photographic plate is put along BD, so that all the rays having a definite value of  $e/m$  fall on the plate at the same point. If rays are present having several values of  $e/m$  a series of lines is produced on the plate, one for each value of  $e/m$ . By varying the electric and magnetic field strengths rays having any desired range of values of  $e/m$  can be brought on to the plate. In this way Aston got well defined lines on his plates, and was able to compare values of  $e/m$  to one part in one thousand by measuring the distances between the lines.

## 7. Aston's Results.

Aston has examined the positive rays from many different elements and finds that the relative masses of the atoms can be expressed by integers within the limits of error. For example, the masses of the rays from carbon, nitrogen, oxygen, and fluorine are proportional to 12, 14, 16, and 19 to within one part in 1000. These numbers agree with the atomic weights 12.00, 14.008, 16, and 19.00. Hydrogen rays form an exception to this rule, for, taking  $O = 16$ , hydrogen rays give  $H = 1.0077$ , in agreement with the chemical atomic weight.

The rays of elements which have atomic weights differing appreciably from integers are found to contain atoms having two or more different masses proportional to integers. For example, chlorine gives rays with masses proportional to 35 and 37, taking oxygen rays proportional to 16 as the standard. The atomic weight of chlorine is 35.46, so that it appears that chlorine is a mixture of two sorts of chlorine atoms having atomic weights 35 and 37. Xenon, the atomic weight of which is 130.2, is found to consist of a mixture of atoms of atomic weights 128, 129, 130, 131, 132, 134, and 136. In this way about 16 elements have been found to consist of mixtures of atoms of different atomic weights.

### 8. Hot Anode Method for Positive Rays.

The discharge tube method of obtaining positive rays is only successful in the case of elements which form stable gaseous compounds which can be admitted to the tube. Positive rays from many very slightly volatile substances can be obtained by means of a hot anode on which a small quantity of the substance is placed. The hot anode is usually a narrow strip of platinum foil which can be heated by passing a current through it. The substance or mixture to be investigated is put on the strip of foil and fused on to it by heating the platinum. The hot anode is mounted directly in front of the cathode of the mass spectrograph and a very good vacuum is maintained in the bulb. When a large potential difference is maintained between the anode and the cathode and the anode is heated, positive rays are emitted by the anode and some of them pass through the slit in the cathode.

The ratio  $e/m$  for alkali metal atoms emitted by alkali sulphates on such a hot anode was determined by O. W. Richardson by measuring their deflections in an electric and transverse magnetic field. Richardson found values of  $e/m$  agreeing closely with those calculated for single atoms which have lost one electron. His experiments were not sufficiently accurate to distinguish between atomic weights differing by only a few units.

Aston and G. P. Thomson examined the positive rays of lithium by the parabola method and got two parabolas corresponding to atomic weights 6 and 7. Aston examined the other alkali metals, using a hot anode and his mass spectrograph. He found sodium to give rays of atomic weight 23 only. Potassium gave rays with atomic weights 39 and 41. Rubidium gave rays of weights 85 and 87, and caesium gave only rays of atomic weight 133.

### 9. Dempster's Method.

The positive rays from several elements have been studied by Dempster by another method. In his apparatus positive rays from a hot anode are accelerated by a known potential difference and then pass through a slit. Immediately beyond the slit they enter a uniform magnetic field which causes them to move along a semi-circular path into another slit. The rays passing through the second slit are received on an insulated electrode and the charge they carry is measured.

If  $P$  is the potential difference used, then

$$Pe = \frac{1}{2}mv^2,$$

where  $e$  is the charge,  $m$  the mass, and  $v$  the velocity of the rays due to

$P$ . If  $H$  is the strength of the magnetic field, and  $r$  the radius of the semicircular path, then

$$\frac{mv^2}{r} = Hev.$$

Hence

$$\frac{m}{e} = \frac{H^2 r^2}{2P}.$$

Dempster kept  $H$  constant, and measured the current received by the insulated electrode as  $P$  was increased by small steps. It was found that the current increased to a maximum at certain values of  $P$ , and the corresponding values of  $m/e$  were calculated. In this way Dempster found magnesium to consist of a mixture of three sorts of atoms having atomic weights 24, 25, and 26. He also found zinc to consist of atoms with weights 63, 65, 67, and 69.

### 10. Isotopes.

It was formerly supposed that all the atoms of an element are precisely equal in all respects, but the methods used to determine atomic weights depended on the relative weights of quantities which contained enormous numbers of atoms. The chemical atomic weights are therefore merely the average atomic weights for very large numbers of atoms. The methods of positive ray analysis first made it possible to test the question whether all the atoms of an element are equal or not.

It is not possible to separate the two sorts of chlorine atoms with atomic weights 35 and 37 by any chemical process. They have precisely similar chemical properties and are therefore regarded as atoms of the same element. The same is true of the different sorts of atoms of other elements. Substances having the same chemical properties but different atomic weights are called *isotopes*.

In a few cases it has been found possible by physical processes such as diffusion or evaporation to slightly alter the relative proportions of the isotopes in certain elements. In this way a quantity of the element is separated into parts having slightly different atomic weights. This was done for neon by Aston, for chlorine by Harkins, and for mercury by Bronsted and Hevesy. The last named physicists prepared two samples of mercury having relative densities 0.99974 and 1.00023 compared with ordinary mercury taken as unity.

The theory of isotopes is discussed in the chapter on Constitution of Atoms. The important application of positive ray analysis to ionization by collisions of electrons, by Smyth, is discussed in the chapter on the Critical Potentials of Atoms.

### REFERENCES

1. *Rays of Positive Electricity*. J. J. Thomson.
2. *Isotopes*. F. W. Aston.

## CHAPTER XI

# Radioactive Transformations

### 1. Discovery of Radioactivity. Radium.

Radioactivity was discovered by Becquerel in 1896. Röntgen's discovery in 1895 of rays which can penetrate materials opaque to ordinary light, and which were apparently associated in some way with the fluorescence of the glass of the Crookes tubes from which they were emitted, suggested the possibility that phosphorescent or fluorescent substances might emit such rays. Becquerel placed a phosphorescent compound of uranium—the double sulphate of uranium and potassium—on a photographic plate wrapped in opaque black paper. He found that the plate was affected and that rays were emitted by the uranium compound which penetrated the paper and other substances. By putting metallic objects between the uranium and the plate shadows of the objects were obtained on the plate.

It was soon found that all compounds of uranium, whether phosphorescent or not, emit the rays, and that the activity depends on the amount of uranium present in the compound and not at all on the other elements present. Many other substances were then tested in the same way. Mme Curie in 1898 found that thorium is also radioactive, and that the mineral pitch-blende and other uranium minerals are four or five times more active than would be expected from the amount of uranium in them. This suggested that uranium minerals contain another radioactive body more active than uranium, and Curie found as the result of a laborious chemical separation of large quantities of pitch-blende residues that there is present in these a substance which has chemical properties similar to barium and which is enormously more radioactive than uranium. This substance was called radium. Another radioactive body, similar to bismuth, was also discovered, and was called polonium.

Curie found that there is about 0.35 gm. of radium to 1 ton of uranium in uranium minerals. Radium is more than one million times more radioactive than uranium. It is not easily separated from barium. Radium bromide is less soluble in water than barium

bromide, which enables them to be separated by a series of crystallizations.

Radium is found to have all the properties of a metallic element. Its atomic weight is 226.0 and it occupies a place in the periodic table of elements in the same column as calcium, strontium, and barium. It gives a spectrum similar to that of the other alkaline earth metals.

Rutherford in 1899 found that the rays from radioactive bodies are of two kinds, which he called  $\alpha$ - and  $\beta$ -rays. The  $\alpha$ -rays are much less penetrating than the  $\beta$ -rays. Villard in 1900 discovered that a third type of rays more penetrating than the  $\beta$ -rays is also emitted, which were called  $\gamma$ -rays. The properties of these three kinds of rays are discussed in other chapters, and it will suffice here to say that  $\alpha$ -rays are positively charged helium atoms shot out with velocities about  $\frac{1}{10}$  that of light,  $\beta$ -rays are high-velocity electrons, and  $\gamma$ -rays are electromagnetic waves of extremely short wave-length,

## 2. Uranium-X and Thorium-X.

An important discovery was made by Sir William Crookes in 1900. He found that if ammonium carbonate is added to a solution of uranium nitrate until the precipitate first formed is redissolved, a small quantity of insoluble precipitate remains which can be filtered off. This small precipitate Crookes found to be strongly radioactive when tested by its action on a photographic plate, whereas the uranium solution is no longer active. The radioactivity of the uranium is therefore due to the presence of a small quantity of another substance, intensely active, which was called uranium-X. It was found later that the uranium-X emits only  $\beta$ -rays, and that the uranium emits  $\alpha$ -rays.  $\alpha$ -rays have not much action on a photographic plate as compared with  $\beta$ -rays, which explains Crookes' results.

The activity of the uranium-X was soon found not to be permanent but to die away gradually so that after a few months it disappeared, whereas the  $\beta$ -ray activity of the uranium gradually recovered after the separation of the uranium-X and after a few months was completely restored. A second quantity of the uranium-X could then be separated from the uranium. It appears therefore that uranium continually produces uranium-X, the activity of which gradually dies away.

Rutherford in 1900 discovered that thorium emits a radioactive gas which was called thorium emanation. The radioactivity of this gas was found to die away in a few minutes. In 1902 Rutherford and Soddy separated an intensely radioactive body from thorium similar to uranium-X, which was called thorium-X. It was found that thorium-X emits the thorium emanation.

M. and Mme Curie discovered that radium also emits an emanation,

and that solid bodies exposed to this emanation become radioactive. The emanation deposits an active material on bodies in contact with it. Rutherford shortly afterwards found that thorium emanation also gives a radioactive deposit. The activity of these active deposits eventually disappears.

### 3. Rutherford and Soddy's Theory.

A theory of radioactivity, now universally accepted, was put forward by Rutherford and Soddy in 1902 to explain the results they had obtained with thorium and its products. According to this theory the atoms of radioactive elements are unstable and eventually explode or decompose into new atoms having quite different properties. The chance that a particular atom will decompose has a definite value, so that if we consider a large number of atoms the number which decompose per unit time is proportional to the number existing. Thus if there are  $N$  atoms at time  $t$ , then

$$\frac{dN}{dt} = -\alpha N,$$

where  $\alpha$  is a constant, exactly as for a monomolecular chemical decomposition. Integrating this equation we get

$$N = N_0 e^{-\alpha t},$$

where  $N_0$  is the number of atoms existing at the time  $t = 0$ . It was found that thorium-X and thorium emanation disappear gradually in accordance with this equation. When  $(N/N_0) = \frac{1}{2}$  we have  $t = (\log_e 2)/\alpha$ , where  $t$  is the time in which  $N$  falls to  $N/2$ . The rate of disappearance of radioactive bodies is conveniently expressed by giving the value of  $(\log_e 2)/\alpha$ , i.e. the time in which the amount remaining drops to one-half the initial value. For thorium emanation this time is 54 sec., and for thorium-X it is 3.64 days. For uranium-X the half value period is 24.6 days.

When a radioactive atom decomposes, the products formed are an  $\alpha$ -ray or a  $\beta$ -ray, but not both, and a new atom. When an  $\alpha$ -ray is one of the products the new atom has an atomic weight four units less than that of the parent atom, because the  $\alpha$ -ray is a positively charged helium atom of atomic weight 4. When a  $\beta$ -ray is emitted the atomic weight of the new atom is almost exactly equal to that of the parent atom, since the mass of a  $\beta$ -ray or electron is less than one thousandth part of that of one hydrogen atom.

According to the nucleus theory of atoms an atom consists of a positively charged nucleus surrounded by electrons. It is supposed that it is the nucleus which decomposes in radioactive changes. Thus when an  $\alpha$ -ray is emitted the charge on the nucleus is diminished by  $2e$ , the charge on the  $\alpha$ -ray, and when a  $\beta$ -ray is emitted the nuclear charge is increased by  $e$ , since  $-e$  is the charge on one electron. Thus when an  $\alpha$ -ray is emitted the atom must also lose two electrons if it remains neutral, and when a  $\beta$ -ray is emitted it must gain one more electron to remain neutral.

Let us suppose that at time  $t = 0$  we have  $N$  atoms of a radioactive substance and that these atoms decompose, giving a second radioactive substance, which decomposes giving a third substance, and so on until finally a non-radioactive product is produced. Let  $N_1, N_2, N_3, \dots, N_n$ , be the numbers of the atoms of the successive products existing at time  $t$ ,  $N_n$  being the number of atoms of the final product.

Then we have

$$\begin{aligned}\frac{dN_1}{dt} &= -\alpha_1 N_1, \\ \frac{dN_2}{dt} &= \alpha_1 N_1 - \alpha_2 N_2, \\ \frac{dN_3}{dt} &= \alpha_2 N_2 - \alpha_3 N_3, \\ &\vdots \\ \frac{dN_n}{dt} &= \alpha_{n-1} N_{n-1}.\end{aligned}$$

The sum of all the terms on the left-hand sides is zero, showing that the total number of atoms remains constant, which is obviously correct. The first equation gives  $N_1 = N e^{-\alpha_1 t}$  since at  $t = 0$ ,  $N_1 = N$ . The solution of the second equation is

$$N_2 = A e^{-\alpha_1 t} + B e^{-\alpha_2 t},$$

where  $A$  and  $B$  are constants. When  $t = 0$ ,  $N_2 = 0$ , so that  $B = -A$  and therefore

$$N_2 = A (e^{-\alpha_1 t} - e^{-\alpha_2 t}).$$

Hence

$$\frac{dN_2}{dt} = -\alpha_1 A e^{-\alpha_1 t} + \alpha_2 A e^{-\alpha_2 t}.$$

When  $t = 0$ , we have  $N_1 = N$  and  $N_2 = 0$ , so that, when  $t = 0$ ,

$$\frac{dN_2}{dt} = \alpha_1 N = A (\alpha_2 - \alpha_1),$$

and therefore

$$A = \frac{\alpha_1 N}{\alpha_2 - \alpha_1}.$$

Hence

$$N_2 = N \frac{\alpha_1}{\alpha_2 - \alpha_1} (e^{-\alpha_1 t} - e^{-\alpha_2 t}).$$

In a similar way we find that

$$N_3 = \frac{N \alpha_1 \alpha_2}{(\alpha_3 - \alpha_1)(\alpha_2 - \alpha_1)(\alpha_3 - \alpha_2)} \{ (\alpha_3 - \alpha_2) e^{-\alpha_1 t} - (\alpha_3 - \alpha_1) e^{-\alpha_2 t} + (\alpha_2 - \alpha_1) e^{-\alpha_3 t} \};$$

and similar expressions may be found for  $N_4$ ,  $N_5$ , &c.

The original substance gradually disappears and the final product gradually increases in amount until  $N_n = N$ . The intermediate products increase up to a maximum amount and then gradually disappear. The time  $t_2$  at which  $N_2$  is a maximum is got by putting  $dN_2/dt = 0$ , which gives

$$t_2 = \frac{1}{\alpha_2 - \alpha_1} \log \frac{\alpha_2}{\alpha_1}.$$

In the same way, the time  $t_3$  at which  $N_3$  is a maximum is found to be given by

$$(\alpha_2 - \alpha_3) \alpha_1 e^{-\alpha_1 t_3} + (\alpha_3 - \alpha_1) \alpha_2 e^{-\alpha_2 t_3} + (\alpha_1 - \alpha_2) \alpha_3 e^{-\alpha_3 t_3} = 0.$$

An important case is that where  $\alpha_1$  is very small compared with the other  $\alpha$ 's. In this case  $N_1$  remains nearly constant and equal approximately to  $N$  for a long time.

The equation  $\frac{dN_2}{dt} = \alpha_1 N_1 - \alpha_2 N_2$ , with  $\alpha_1 N_1$  constant, gives

$$N_2 = N_1 \frac{\alpha_1}{\alpha_2} (1 - e^{-\alpha_2 t}),$$

which shows that  $N_2$  increases to a nearly constant value  $N_1 \frac{\alpha_1}{\alpha_2}$ .

The equation  $\frac{dN_3}{dt} = N_2 \alpha_2 - \alpha_3 N_3$ ,

with  $N_2$  constant and equal to  $N_1 \alpha_1 / \alpha_2$ , becomes

$$\frac{dN_3}{dt} = N_1 \alpha_1 - \alpha_3 N_3,$$

which gives

$$N_3 = N_1 \frac{\alpha_1}{\alpha_3} (1 - e^{-\alpha_3 t}),$$

so that  $N_3$  also increases to a constant value  $N_1 \alpha_1 / \alpha_3$ . In the same way we find that  $N_4$  becomes constant and equal to  $N_1 \alpha_1 / \alpha_4$ , and so on. Hence in this case we find that eventually

$$N_1 \alpha_1 = N_2 \alpha_2 = N_3 \alpha_3 = N_4 \alpha_4 = N_5 \alpha_5 = \dots,$$

provided the  $\alpha$ 's are large compared with  $\alpha_1$ . Since  $\alpha_n = 0$ , we do not get  $N_1 \alpha_1 = N_n \alpha_n$ , and clearly  $\frac{dN_n}{dt} = \alpha_{n-1} N_{n-1} = N_1 \alpha_1$ , so that the final product increases at the same rate that the first body diminishes, while all the intermediate products remain nearly constant in amount. The amounts of the intermediate products are all proportional to  $N_1$  and so diminish very slowly as  $N_1$  diminishes. When the state for which

$$N_1 \alpha_1 = N_2 \alpha_2 = N_3 \alpha_3 = \dots$$

has been attained, the mixture is said to be in a state of radioactive equilibrium, because each intermediate product is produced almost as fast as it decomposes.

For any intermediate product we have

$$\frac{dN_S}{dt} = \alpha_{S-1} N_{S-1} - \alpha_S N_S,$$

where  $S = 2, 3, 4, \dots, n-1$ .

It is clear that if  $N_S$  is constant for any value of  $S$  from 2 to  $n-1$ , then

$$\alpha_{S-1} N_{S-1} = \alpha_S N_S,$$

so that

$$\alpha_1 N_1 = \alpha_2 N_2 = \alpha_3 N_3 = \alpha_4 N_4 = \dots = \alpha_{n-1} N_{n-1}.$$

Since

$$\frac{dN_n}{dt} = \alpha_{n-1} N_{n-1},$$

we see that in this case  $N_n = \alpha_1 N_1 t + \text{constant}$ , so that the amount of the final product increases at a uniform rate. We have also  $\alpha_1 N_1 = \text{constant}$ , so that in order that the intermediate products should remain constant  $N_1$  must be supposed kept constant in some way. This condition is approximately satisfied when  $\alpha_1$  is very small. The product  $\alpha_S N_S$  is equal to the number of atoms of the body of number  $S$  which decompose in unit time, so that, in a mixture in radioactive equilibrium, the number of atoms decomposing is the same for all the intermediate products.

The number of  $\alpha$ -rays emitted by any one of the products present is therefore

the same as the number emitted by any other product which emits  $\alpha$ -rays. The same is true of the emission of  $\beta$ -rays by the atomic nuclei. The amounts of the different bodies present in the mixture are inversely as their activities, so that the bodies which decompose rapidly are present in small amounts. Uranium minerals are examples of such a mixture of radioactive bodies in a state of radioactive equilibrium.

#### 4. Table of Products formed from Uranium.

The following table gives the products formed from uranium. Each product is formed by the decomposition of the atoms of the preceding product in the table. The first column gives the names and the second the chemical symbols of the products. The third column gives the atomic weights. These are got from that of uranium by subtracting 4 when  $\alpha$ -rays are emitted. The next column gives the atomic number. This increases by one when  $\beta$ -rays are emitted and diminishes by two when  $\alpha$ -rays are emitted. The atomic weight of radium has been found directly by chemical analysis. The fifth column gives the symbol of an isotope of the product, that is, an element which has the same atomic number and identical chemical properties. The chemical properties of radium are very similar to those of barium, and radon is a chemically inactive gas like argon. Its atomic weight has been determined by measuring its density. The sixth column headed T gives the half-value period. The next column gives the rays emitted. The last column gives the velocity of the rays as a fraction of that of light. Rays in brackets are relatively feeble.

Two other similar series of radioactive products are known, one starting with thorium and ending with lead, and the other of uncertain origin containing actinium.

Name.	Symbol.	Atomic Weight.	Atomic Number.	Iso- tope.	T.	Radia- tion.	Velocity of Rays.
Uranium I ...	U-I	238	92	—	$4.67 \times 10^9$ yr.	$\alpha$	0.0456
Uranium $X_1$ ...	U- $X_1$	234	90	Th	24.6 days	$\beta$	—
Uranium $X_2$ ...	U- $X_2$	234	91	Pa	1.15 min.	$\beta(\gamma)$	—
Uranium II ...	U <sub>II</sub>	234	92	U	$2 \times 10^6$ yr.	$\alpha$	0.0479
Ionium ...	Io	230	90	Th	$6.9 \times 10^4$ yr.	$\alpha$	0.0485
Radium ...	Ra	226	88	—	1690 yr.	$\alpha(\beta, \gamma)$	0.050
Ra Emanation } (Radon) ... }	Rn	222	86	—	3.85 days	$\alpha$	0.054
Radium A ...	Ra-A	218	84	Po	3.0 min.	$\alpha$	0.0565
Radium B ...	Ra-B	214	82	Pb	26.8 min.	$\beta(\gamma)$ {	0.36; 0.41; 0.63; 0.70; 0.74
Radium C ...	Ra-C	214	83	Bi	19.5 min.	$\beta$ {	0.786; 0.862; 0.949; 0.957
Radium C' ...	Ra-C'	214	84	Po	$10^{-6}$ sec.	$\alpha$	0.0641
Radium D ...	Ra-D	210	82	Pb	16.5 yr.	$(\beta, \gamma)$	0.33; 0.39
Radium E ...	Ra-E	210	83	Bi	5 days	$\beta$	—
Radium F } (Polonium) }	Ra-F	210	84	Po	136 days	$\alpha(\gamma)$	0.0523
Radium $\Omega$ } (Lead) ... }	Ra- $\Omega$	206	82	Pb	—	—	—

Radium-C also gives another product to the extent of 0.03 per cent which is produced with the emission of  $\alpha$ -rays. This product is called radium-C''; its atomic weight is 210, its atomic number 81; its isotope thallium and its half-value period 1.4 min. It emits only  $\beta$ -rays. Radium apparently emits both  $\alpha$ - and  $\beta$ -rays, which is unusual. The  $\beta$ -rays are feeble, and may be due to a few radium atoms decomposing in a different way from that which gives the emanation. If a radium atom emitted an  $\alpha$ -ray and also a  $\beta$ -ray the resulting atom should have atomic weight 222 and atomic number 87. An element with atomic number 87 should have chemical properties similar to those of caesium, whereas the emanation is chemically inert, as it should be if its atomic number is 86.

### 5. Properties of Radon.

It is only possible here to give an account of the investigations which have been made of the properties of a very few of the radioactive bodies. An account will be given of radium emanation or radon, and some of its products. Radon is produced from radium at a practically constant rate, since the half-value period of radium is 1690 years. The radon atoms do not escape to any great extent from solid radium salts since they are produced throughout the volume of the solid and do not diffuse appreciably through it. The radon therefore accumulates in solid radium salts until its rate of decomposition is equal to its rate of production. If the salt is dissolved in water the radon is able to escape, especially if the water is heated under a low pressure.

For the purpose of obtaining a supply of radon it is convenient to keep a solution of some radium salt in a glass flask connected to a Toepler pump, by means of which the gases evolved by the solution may be pumped off and collected over mercury. It is found that the solution evolves small amounts of hydrogen and oxygen, which are evidently produced by the action of the radiations on the water. The radon is therefore obtained mixed with oxygen, hydrogen, and water vapour. If all the radon in the solution is collected in this way then after a few days another supply will have accumulated.

The radon may be obtained in the pure state by means of the apparatus shown in fig. 1. The mixture of radon and other gases over mercury in the tube A is let into a bulb B through the stopcock C, the bulb B having been previously exhausted through E, which leads

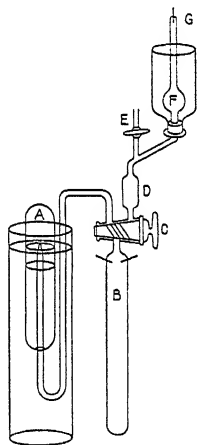


Fig. 1

to a Toepler pump. The mixture is exploded in B by means of a spark, and  $\text{CO}_2$  is absorbed by means of some caustic potash fused on to the walls of B. The gases are passed into a bulb F through D, which contains phosphorus pentoxide to absorb the water vapour. The bulb F is then surrounded by liquid air and the radon condenses on its walls. The apparatus is next pumped out through E and mercury let in so as to fill it up to the bottom of F. On the liquid air being removed the emanation evaporates and fills F. It can be compressed into a small calibrated capillary tube G at the top of the bulb F by letting in more mercury. In this way the volume of the emanation at different pressures can be measured.

Ramsay and Soddy obtained pure radon from a solution containing 60 mgm. of radium in this way. They found the gas obeyed Boyle's law, and that its volume gradually decreased, falling to half-value in about 4 days. The  $\alpha$ -rays penetrate the glass walls and so may not produce gaseous helium in the tube, otherwise the volume should not have diminished. Later experiments have shown that the volume of radon in equilibrium with 1 gm. of radium is approximately 0.6 c. mm. This quantity of radon is called a *curie*.

The number of  $\alpha$ -rays emitted by radium, free from other products, has been counted and found to be  $3.4 \times 10^{10}$  per second from 1 gm. of radium. This should be therefore the number of atoms of radon produced per second. The rate at which the atoms of radon decompose is  $\alpha N$ , where  $\alpha = 2.085 \times 10^{-6}$  and  $N$  is the number of atoms of radon. Thus if  $3.4 \times 10^{10}$  new atoms are produced per second the number existing when equilibrium is reached will be

$$N = \frac{3.4 \times 10^{10}}{2.085 \times 10^{-6}} = 1.63 \times 10^{16}.$$

The number of atoms in 1 c. c. of a monatomic gas at  $0^\circ \text{C}$ . and 760 mm. is  $2.7 \times 10^{19}$ , so that the volume of radon in equilibrium with 1 gm. of radium should be

$$\frac{1.63 \times 10^{16}}{2.7 \times 10^{19}} \times 1000 = 0.603 \text{ c. mm.},$$

which agrees very well with the value found experimentally.

The rate of decomposition of radon has been accurately determined by Curie and by Rutherford. Some radon was sealed up in a glass tube, and the intensities of the  $\beta$  and  $\gamma$  radiations coming out of the tube compared at measured times by means of the ionization produced by these rays in a metal ionization chamber. The radon itself emits only  $\alpha$ -rays, but the product radium-C gives  $\beta$ - and  $\gamma$ -rays. The half-value periods of radium-A, radium-B, and radium-C are all small

compared with that of radon, so that in a short time the amount of radium-C present in the tube becomes practically proportional to the amount of radon present, and then the rate of decay of the  $\beta$ - and  $\gamma$ -rays is a measure of the decay of the radon. In this way it was found that  $\alpha$  for radon is  $2.085 \times 10^{-6}$  (sec.) $^{-1}$ , which gives 3.85 days for the half-value period.

Radon is found to be without chemical properties, like the inert gases helium, neon, argon, krypton, and xenon. Its density was determined by Ramsay and Gray by weighing a fraction of a cubic millimetre of it in a bulb with a microbalance of extraordinary sensibility. The density was nearly 222, taking that of oxygen to be 16, and so agreed with the atomic weight got by deducting four from 226, the atomic weight of radium. Radon therefore comes at the end of the sixth period of the periodic arrangement of the elements, in the same column as the other inert gases. The atomic number of radon is 86, which is got from that of radium 88 by deducting 2 for the emission of one  $\alpha$ -ray.

The atomic numbers of the inert gases 2, 10, 18, 36, 54, 86 are equal respectively to  $2 \times 1$ ,  $2(1 + 2^2)$ ,  $2(1 + 2^2 + 2^2)$ ,  $2(1 + 2^2 + 2^2 + 3^2)$ ,  $2(1 + 2^2 + 2^2 + 3^2 + 3^2)$ , and  $2(1 + 2^2 + 2^2 + 3^2 + 3^2 + 4^2)$ , and so form a regular series with that of radon as the last term.

## 6. Experimental Study of Products from Radon.

Solid bodies put in contact with radon become coated with a radioactive deposit. This deposit can be concentrated on to a small wire by charging the wire negatively to a few hundred volts. Thus if a solution of radium bromide is put at the bottom of a corked bottle, and a fine wire put through the cork and kept negatively charged, the wire becomes strongly radioactive in a short time. The radiations from the wire may be studied by means of the ionization which they produce in air.

An electroscope suitable for the study of the ionization due to  $\alpha$ -rays is shown in fig. 2. It consists of a cubical metal box A containing two parallel metal plates B and C. The plate B is insulated and connected to a rod D which supports a narrow gold leaf E. The gold leaf is in an upper chamber above A. The upper end of D is covered by a removable cap at G which enables the rod to be charged. The deflection of the leaf can be observed with a low-power microscope through the glass window F. A graduated scale in the eyepiece of the microscope enables the deflections to be measured. If a body emitting

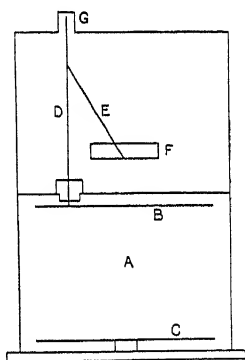


Fig. 2

$\alpha$ -rays is put on the lower plate C the rays ionize the air and the leaf gradually falls. The number of scale divisions moved by the leaf in a suitable interval of time is proportional to the ionization. To study the  $\beta$ -rays emitted by any radioactive body it is enclosed in a tube or covered with a thin sheet thick enough to stop the  $\alpha$ -rays. If it is then put on the plate C the rate of fall of the leaf measures the intensity of the ionization due to the  $\beta$ - and  $\gamma$ -rays. Ionizations due to  $\gamma$ -rays alone may be studied by absorbing the  $\alpha$ - and  $\beta$ -rays by means of thick metal plates.

If a bare wire covered with the deposit from radon is put in such an electroscope the ionization is almost entirely due to the  $\alpha$ -rays. If the wire has only been exposed to the radon for a short time, say 1 min., the deposit will consist almost entirely of the product of the decomposition of the radon which is called radium-A. It is found that the intensity of the  $\alpha$ -rays decreases rapidly for about 10 min., the half-value period being 3 min. It then becomes nearly constant for about 1 hr. and then begins to fall again, with a half-value period of 28 min. The  $\beta$ -ray activity of a wire which has been exposed to radon for less than 1 min. is small initially, but rises in about 20 min. to a maximum and then decreases. This shows that radium-A gives no  $\beta$ -rays, but forms products which give them. The variation of the  $\alpha$ -ray activity shows that the products also give some  $\alpha$ -rays. On heating the wire to about 600° C. it is found that part of the deposit volatilizes and part remains on the wire. The part remaining on the wire gives  $\alpha$ - and  $\beta$ -rays, which both decrease with the half-value period of 19.5 min. The part which evaporates can be condensed on a cool surface. It is inactive at first but then emits  $\alpha$ - and  $\beta$ -rays, which increase in intensity to a maximum and then die away.

The results obtained can be explained satisfactorily as follows. The radium-A emits  $\alpha$ -rays only and has a half-value period of 3 min. It gives radium-B with a half-value period of 26.8 min., which emits only very feeble  $\beta$ -rays. The radium-B gives radium-C, which emits  $\beta$ -rays and has a half-value period of 19.5 min. This gives radium-C' with a very short period which gives  $\alpha$ -rays. The  $\alpha$ -rays from radium-C' and the  $\beta$ -rays from radium-C both vary with the time in the same way because of the very short period of radium-C'.

It is found that the radioactivity of the deposit does not entirely disappear for a long time, and it appears that several more products are formed. The next one, radium-D, has a half-value period of 16.5 years.

## 7. Heat evolved by Radium.

When the rays from radioactive bodies are absorbed by matter the energy of the rays is nearly all converted into heat. It is found, for example, that 1 gm. of radium in equilibrium with its products pro-

duces about 120 calories per hour. The total heat produced by a gramme of radium is therefore

$$\frac{120}{60 \times 60} \int_0^{\infty} e^{-at} dt, \text{ where } a = 1.3 \times 10^{-11} \text{ sec.}^{-1},$$

which is the value of  $a$  corresponding to a half period of 1690 years. This is equal to  $2.57 \times 10^9$  calories. This is enormously greater than the heat evolved per gramme in any chemical reaction. The source of this energy will be discussed in the chapter on the Constitution of Atoms.

#### REFERENCE

*Radiactivity.* E. Rutherford.

## CHAPTER XII

### Constitution of Atoms

#### 1. Atomicity of Electricity and of Matter. Electrons and Protons.

In this chapter it is proposed to discuss the modern theory of atoms, which is supported by many of the facts described in more or less detail in the preceding chapters.

Many of the quantities which are measured experimentally usually are, or at any rate appear to be, continuous, that is they are capable of having any value between certain limits, such quantities as length, mass, and time for example. A quantity is said to be atomic, or made up of atoms, when its values are always integral multiples of a definite value, which is the smallest value of the quantity, not zero, which occurs. Thus, as we have seen, when a very small quantity of electricity is measured, it is found to be equal to  $4.77 \times 10^{-10}$  electrostatic units, multiplied by an integer. Electricity is therefore regarded as being made up of atoms of electricity, the negative electrons and the protons.

By means of Aston's mass spectrograph it has been shown that the ratio of the mass ( $m$ ) to the charge ( $E$ ) of very small particles charged with positive electricity is always an integral multiple of a definite value. That is  $m/E = Nf$ , where  $N$  is an integer and  $f$  is the smallest value of  $m/E$  observed. But  $E = ne$ , where  $n$  is an integer and  $e$  is the charge of the atom of electricity, so that we must have  $m = nNef$ . The mass  $m$  is therefore equal to  $ef$  multiplied by  $nN$ , which is an integer.

It appears, therefore, that matter is atomic, the atomic mass being equal to  $ef$ , the value of  $m$  when  $nN = 1$ . Now  $f$  is found to be equal to  $1/9650$  with the charge expressed in electromagnetic units, so that  $ef = (4.77 \times 10^{-10}) / (3 \times 10^{10} \times 9650) = 1.66 \times 10^{-24}$  gm. Matter is therefore made up of atoms each of which has a mass  $1.66 \times 10^{-24}$  gm. and a charge  $4.77 \times 10^{-10}$  electrostatic units.

These atoms, of course, are not the atoms of the chemist, which must be made up out of them. These ultimate atoms of matter shown to exist by means of the mass spectrograph are called protons. The chemical atomic weights which are shown by the mass spectrograph

to be integers when that of oxygen is, as usual, taken equal to 16, are equal to the number of protons in one atom of the different elements.

An important exception occurs in the case of hydrogen, the atomic weight of which is 1.007 and so not exactly an integer. This is believed to be due to energy escaping when two protons combine, so that a single proton has slightly more mass than the average mass of the protons combined together in an atom. The values found for  $m/E$  are integral multiples of  $1/9650$ , but the smallest value of  $m/E$  observed, which is got with hydrogen, is  $1.007/9650$ .

As we have seen, negative electrons are also contained in all kinds of matter.  $e/m$  for electrons is equal to  $1.76 \times 10^7$  electromagnetic units per gramme. There are good reasons for believing that the charge on one electron is equal to that of the atom of electricity or  $4.77 \times 10^{-10}$  electrostatic units. It is found, for example, that when the oil droplet in Millikan's apparatus for measuring small charges is illuminated by ultra-violet light its charge is increased by that of one atom of electricity at a time, and it is known that  $e/m$  for the charges set free by ultra-violet light in a vacuum is the same as for electrons. There is no reason to suppose that the charges liberated from bodies by ultra-violet light are not the same in a vacuum as in air at atmospheric pressure, so that the electronic charge must be just one atom of electricity. The mass of one electron is therefore equal to  $(4.77 \times 10^{-10}) / (3 \times 10^{10} \times 1.76 \times 10^7) = 9 \times 10^{-28}$  gm.

Owing to the small mass of electrons the number of them in an atom makes very little difference to the atomic weight. It is supposed therefore that matter consists of protons and electrons, and that the charge on one proton is  $+4.77 \times 10^{-10}$  and that on one electron  $-4.77 \times 10^{-10}$ . The total number of electrons in any electrically neutral portion of matter must therefore be equal to the number of protons. In particular any neutral atom must contain as many electrons as protons.

## 2. Nucleus Theory of Atoms. Atomic Number.

Rutherford's experiments on the scattering of  $\alpha$ -rays strongly support the theory that atoms consist of a positively charged nucleus with electrons around it. This theory is also supported by Bohr's quantum theory of optical and X-ray spectra. The charge on the nucleus must be a multiple of the atom of electricity or protonic charge which will be denoted as usual by  $e$ . We should therefore expect to find atoms with nuclear charges  $e, 2e, 3e, 4e \dots$ . It is natural to expect the nuclear charge to increase with the atomic weight, and it is supposed that the nuclear charge is equal to  $Ne$ , where  $N$  is the atomic number or number giving the position of an element in a list of the elements arranged in order of increasing atomic weights. Thus  $N$  for hydrogen is one, for helium two, for lithium three, and so on. This

idea is strongly supported by Moseley's discovery that the square root of the frequency of the  $K$  X-rays of any element is a linear function of the atomic number  $N$ . This fundamentally important discovery shows that some quantity in atoms increases by the same amount in passing from any atom to the atom of next higher atomic weight. It is difficult to think of any other quantity besides the nuclear charge which could increase in this way. The values of the nuclear charge obtained from the observed scattering of  $\alpha$ -rays by different elements agree closely with  $N$ , so that it is now generally agreed that this is the value of the nuclear charge. It follows that the number of electrons around the nucleus in an electrically neutral atom is equal to  $N$ .

Since a neutral atom of atomic weight  $A$  contains  $A$  protons and  $A$  electrons and there are  $N$  electrons outside the nucleus, the number of electrons inside the nucleus must be  $A - N$ . Thus the hydrogen atom consists of one proton with one electron outside; the helium atom of four protons with two electrons inside the nucleus and two outside. A lithium atom of atomic weight 7 has three electrons outside the nucleus and four inside, while a lithium atom of atomic weight 6 has three electrons outside the nucleus and three inside. The atomic weight is never less than twice the atomic number, so that at least half the electrons are always inside the nucleus, except in the case of the hydrogen atom.

### 3. Electronic Arrangement and Chemical Properties.

It is supposed that the properties of atoms are determined by the number of electrons outside the nucleus. The existence of isotopes, or atoms having practically identical properties but different atomic weights, is therefore explained by supposing that all the isotopes of an element have the same nuclear charge, or the same number of electrons outside the nucleus when the atoms are electrically neutral.

It is found that the properties of atoms vary periodically with the atomic number. Thus the atoms having the atomic numbers 2, 10, 18, 36, 54, 86 are all inert gases having no chemical properties. It appears that these numbers of electrons form stable systems which do not attract other atoms. They must excite relatively weak fields in the space around them.

The atoms having one electron less than the inert gases, those with atomic numbers 9, 17, 35, 53, are all strongly electronegative monovalent non-metallic elements, that is, these atoms strongly attract an electron and form stable compounds with atoms which easily lose an electron.

The atoms which have one more electron than the inert gases, with atomic numbers 3, 11, 19, 37, 55, are all strongly electropositive alkali metals. They combine with the elements having atomic numbers 9, 17, 35, 53, giving very stable compounds. It is supposed that these

atoms easily lose one electron, and that an atom which easily loses one electron combines readily with one which strongly attracts an electron. This may be explained by supposing that the numbers of electrons in the inert gases form a stable system, so that an atom with one more electron than an inert gas atom and another with one less combine, giving a molecule in which there are two stable sets of electrons, each like the stable set in an inert gas atom. The atom which gains an electron is negatively charged, and the other one positively charged, so that they attract each other.

Many of the chemical properties of atoms can be explained qualitatively by this theory, which was first put forward by J. J. Thomson and has been elaborated by Langmuir and others.

As another example consider the chemical radical  $\text{NH}_4$ , which, as is well known, has chemical properties very similar to those of the alkali metals. This radical contains eleven electrons outside the nuclei, which is the same number as in the sodium atom. On losing one electron ten remain, the number which forms the stable system of the neon atom. Again, methane ( $\text{CH}_4$ ) is a very stable compound which does not easily combine with other bodies; its molecules have ten electrons outside the nuclei, like the inert gas neon.

An atom of atomic number  $N$  between those of two inert gases, for example neon (10) and argon (18), can form a stable set of electrons either by losing electrons till ten are left or by gaining enough electrons to make up eighteen. The atom may therefore behave as an electronegative atom of valency  $18 - N$  or as an electropositive atom of valency  $N - 10$ . The sum of the two possible valencies is  $18 - N + N - 10 = 8$ . For example, chlorine with  $N = 17$  is monovalent in salts like  $\text{NaCl}$  but heptavalent in  $\text{KClO}_3$ .

#### 4. The Periodic Table.

On p. 210 is a list of the elements, giving the atomic numbers, average so-called chemical atomic weights, and the actual atomic weights as found with the mass spectrograph.

The elements in the middles of the fourth, fifth, and sixth periods have been omitted. The seventh period is incomplete since no elements are known with atomic numbers greater than 92.

#### 5. Periodicity and the Grouping of Electrons.

To explain the periodic variation of the properties of atoms with the atomic number  $N$ , it is supposed that the electrons move in groups of orbits round the nucleus. The chemical properties are supposed to depend largely on the number of electrons in the outermost group, so that elements like fluorine, chlorine, bromine, and iodine which have very similar properties are supposed to have equal numbers of electrons in the outer group.

Element.		Symbol.	Atomic Number.	Principal Valency.	Chemical Atomic Weight.	Actual Atomic Weights.
First Period.	Hydrogen	H	1	1+	1.008	1.008
	Helium ..	He	2	0	4.0	4
Second Period.	Lithium ..	Li	3	1+	6.94	6, 7
	Beryllium	Be	4	2+	9.1	9
	Boron ..	B	5	3	10.9	10, 11
	Carbon ..	C	6	4	12.00	12
	Nitrogen ..	N	7	3—	14.008	14
	Oxygen ..	O	8	2—	16	16
	Fluorine ..	F	9	1—	19.00	19
	Neon ..	Ne	10	0	20.20	20, 22
Third Period.	Sodium ..	Na	11	1+	23.00	23
	Magnesium	Mg	12	2+	24.32	24, 25, 26
	Aluminium	Al	13	3+	26.96	27
	Silicon ..	Si	14	4	28.3	28, 29, 30
	Phosphorus	P	15	3	31.04	31
	Sulphur ..	S	16	2—	32.06	32, 33, 34
	Chlorine ..	Cl	17	1—	35.46	35, 37, 39
	Argon ..	A	18	0	39.9	36, 40
Fourth Period.	Potassium	K	19	1+	39.1	39, 41
	Calcium ..	Ca	20	2+	40.07	40, 44
	.....					
	Selenium ..	Se	34	2—	79.2	—
	Bromine ..	Br	35	1—	79.92	79, 81
Fifth Period.	Krypton ..	Kr	36	0	82.92	78, 80, 82, 83, 84, 86
	Rubidium	Rb	37	1+	85.45	85, 87
	Strontium	Sr	38	2+	87.63	—
	.....					
	Iodine ..	I	53	1—	126.92	127
Sixth Period.	Xenon ..	X	54	0	130.2	129, 131, 132, 134, 136
	Cæsium ..	Cs	55	1+	132.81	133
	Barium ..	Ba	56	2+	137.37	—
	.....					
	Radon ..	Rn	86	0	222.0	—
Seventh Period.	—	—	87	—	—	—
	Radium ..	Ra	88	2+	226.0	—
	Actinium ..	Ac	89	—	—	—
	Thorium ..	Th	90	—	232.15	—
	Uranium-X	U-X	91	—	—	—
	Uranium ..	Ur	92	—	238.2	—

The theory of the grouping of the electrons in atoms is mainly due to Bohr. It has recently been improved by Main-Smith and Stoner.

The electrons are supposed to move in groups of orbits, all those

in each group having the same principal quantum number  $n$ . These groups correspond to the  $K, L, M, N, \dots$  groups of energy levels. For the  $K$  level  $n = 1$ , for the  $L$  level  $n = 2$ , and so on. There are 1  $K$  level, 3  $L$  levels, and 5  $M$  levels. The level to which an electron belongs in a group is indicated by the second quantum number  $n_1$  and the inner quantum number  $n_2$ .  $n_1$  is equal to or less than  $n$ , so that since there are  $2n - 1$  levels in a group the third number is required. It is supposed to be equal to  $n_1$  or to  $n_1 - 1$ . This gives  $2n - 1$  levels in the groups in agreement with the number of  $K, L$ , and  $M$  levels observed.

The problem now is to determine the number of electrons in each level in the different atoms. The total number of electrons to be placed is equal to the atomic number of the atom, and it is believed that the maximum number in any level is equal to  $2n_2$ . The grouping of the electrons in the inert gas atoms is supposed to be as follows:

	1			2			3			4				5				6		
"	1	1	2	1	2	3	1	2	3	1	2	3	4	1	2	3	4	1	2	3
" <sub>1</sub>	1	1	1	2	1	1	2	2	3	1	1	2	2	3	3	4	1	1	2	3
" <sub>2</sub>	1	1	1	2	1	1	2	2	3	1	1	2	2	3	3	4	1	1	2	3
(2) Helium ...	2																			
(10) Neon ...	2	2	2	4			-													
(18) Argon ...	2	2	2	4																
(36) Krypton ...	2	2	2	4			2	2	4											
(54) Xenon ...	2	2	2	4			2	2	4	4	6									
(86) Radon ...	2	2	2	4			2	2	4	4	6									

The alkali metals have one more electron than the inert gases, and the additional electron is supposed to be in a group by itself outside the other electrons. Sodium has two electrons in group one, eight in group two, like neon, but in addition one in group three. Thus the alkali metals all have a single electron describing an outer orbit by itself, and so have similar properties.

In the same way the alkaline earth metals are supposed to have their electrons grouped like those of the inert gases, but with two additional electrons in an outer group by themselves.

The elements fluorine, chlorine, bromine, and iodine have their electrons grouped like the inert gases, except that one electron is missing from the outermost group. The elements oxygen, sulphur, selenium, and tellurium have two electrons missing as compared with the inert gases; and so on for the other elements.

## 6. Radioactivity and the Nucleus Theory.

The phenomena of radioactivity indicate that the nuclei of the atoms of atomic number greater than 85 are more or less unstable and ultimately decompose with the emission of  $\alpha$ -rays or electrons. It is

remarkable that protons are never emitted by radioactive bodies. An  $\alpha$ -ray consists of four protons and two electrons and is evidently a very stable system. The emission of  $\alpha$ -rays and not protons suggests that the protons in atomic nuclei are often arranged in groups of four or as  $\alpha$ -ray particles in the nucleus. The nuclei of atoms having atomic weights which are multiples of four may consist entirely of  $\alpha$ -ray particles, in which case the atomic number would be one-half the atomic weight as for helium. Atoms with atomic weights 4, 12, 16, 20, 24, 28, 32, 36, 40 are known with atomic numbers half the atomic weight, but for atomic numbers above twenty the atomic weight is greater than twice the atomic number.

Part at any rate of the kinetic energy of the  $\alpha$ -rays must be due to the repulsion between the nucleus and the ray. If we assume that the charges of the nucleus and of the  $\alpha$ -ray can be regarded as point charges, we can easily calculate the initial distance between them corresponding to the kinetic energy of the ray.

If  $N$  is the atomic number of the atom after the ray has been emitted, and assuming all the energy goes into the  $\alpha$ -ray,

$$\frac{Ne}{r} \times 2e = \frac{1}{2}mv^2.$$

For radon,  $N = 84$  and  $v = 0.054 \times 3 \times 10^{10}$  cm. per sec., which with  $e = 4.77 \times 10^{-10}$  and  $e/m = \frac{1}{4} \times 9650 \times 3 \times 10^{10}$  gives  $r = 4.4 \times 10^{-12}$  cm., which is of the same order as estimates made of the radii of atomic nuclei by other methods.

## 7. Ejection of Protons from the Nucleus.

Rutherford made the very interesting discovery that when high-velocity  $\alpha$ -rays are passed through certain substances then rays are produced which have a greater range than the  $\alpha$ -rays. These rays are found to be protons, and are supposed to be produced by collisions of the  $\alpha$ -rays with atomic nuclei. It appears that the  $\alpha$ -ray enters the nucleus and remains in it and causes the emission of a proton. This must increase the atomic weight of the atom by three and its atomic number by unity. This effect was first obtained with nitrogen.  $\alpha$ -rays from radium-C' were passed through nitrogen and the scintillations they produced on a zinc sulphide screen observed. When the distance between the radium-C' and the screen is gradually increased the scintillations disappear abruptly when the distance becomes greater than the range of the  $\alpha$ -rays. However, it was found that a few scintillations were obtained at distances much greater than this range. These scintillations were feebler than the  $\alpha$ -ray scintillations and resembled those produced by the high-velocity protons obtained when  $\alpha$ -rays are passed through hydrogen.

The magnetic and electric deflections of these rays were deter-

mined, and it was found that they carry a positive charge and that  $e/m$  is equal to the value for protons. It is clear that they are high-velocity protons. It is found that only one high-velocity proton is produced by several hundred thousand  $\alpha$ -rays. By passing the  $\alpha$ -rays through thin sheets similar high-velocity protons have been obtained from several solid elements.

The protons come out in all directions, but more come out in directions near that of the  $\alpha$ -rays than in any other direction. Protons have been obtained in this way from boron, nitrogen, fluorine, sodium, aluminium, and phosphorus, and probably from beryllium, magnesium, and silicon. The atomic numbers, atomic weights, and ranges of the protons emitted are as follows:

Element.	Atomic Number.	Atomic Weight.	Forward Range.	Backward Range.
			Cm.	Cm.
Boron .. ..	5	11, 10	58	38
Nitrogen .. .	7	14	40	18
Fluorine .. .	9	19	65	48
Sodium .. .	11	23	58	36
Aluminium ..	13	27	58	36
Phosphorus ..	15	31	65	49

All these elements have odd atomic numbers and their atomic weights are not multiples of four. Elements like oxygen and carbon, the atoms of which can be regarded as made up of  $\alpha$ -rays, do not give protons when bombarded by  $\alpha$ -rays.

It is found that the kinetic energy of the protons is greater than that of the  $\alpha$ -rays, by about forty per cent in the case of aluminium, so that it appears that the atomic nuclei give up some internal energy to the escaping protons. It has been suggested that  $\alpha$ -ray particles and protons inside the nucleus may be describing orbits round a central charge, just as outside the nucleus the electrons move in orbits round the nucleus. The possible orbits inside the nucleus may be determined by quantum principles similar to those which appear to determine the possible orbits of the electrons outside the nucleus. Owing to the masses of the  $\alpha$ -ray particles and protons being enormously greater than the mass of an electron, their possible orbits on the quantum theory are very much smaller than the orbits of electrons. We may suppose that when an  $\alpha$ -ray particle enters the nucleus of an atom it may cause a proton describing an orbit near the surface of the nucleus to be ejected.

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## CHAPTER XIII

### Gaseous Ions

#### 1. Mobility of Ions.

When gases are made to conduct electricity by means of X-rays, ultra-violet light, or other agencies, it is supposed that the conductivity is due to the presence of minute electrically charged particles which are called ions. The properties of these ions is the subject to be discussed in this chapter.

The velocity of ions due to an electric field depends on the temperature, pressure, and nature of the gas in which they are moving. It is proportional to the strength of the electric field, and the velocity due to a field of unit strength, usually one volt per centimetre, is called the mobility of the ions. A group of a large number of ions in a uniform electric field moves with a definite velocity, that of the centre of mass of the group, but this is not true of each ion in the group. The ions diffuse through the gas when there is no field, and in a field they still diffuse, so that the displacements due to diffusion are added to those due to the motion caused by the electric field. In a strong field the motion due to diffusion may be very small compared with that due to the field. The mobility of a positive ion will be denoted by  $k_1$  and that of a negative ion by  $k_2$ .

If a gas contains  $n_1$  positive ions per unit volume, and  $n_2$  negative ions, the current density in it due to an electric field of strength  $F$  is

$$i = (n_1 e_1 k_1 + n_2 e_2 k_2) F,$$

where  $e_1$  is the charge on a positive ion and  $e_2$  that on a negative ion. The products  $e_1 k_1$  and  $e_2 k_2$  are both positive, because changing the sign of  $e$  also changes the sign of  $k$ . Practically all gaseous ions are found to carry charges of the same magnitude of either positive or negative electricity. This ionic charge is equal to the charge on one hydrogen ion in a solution, or to the charge carried by one electron. It will be denoted by  $e$ . The current density is therefore given by

$$i = (k_1 n_1 + k_2 n_2) e F,$$

where both  $k_1$  and  $k_2$ , as well as  $e$ , are now regarded as positive.

## 2. Measurement of Ionic Mobilities. Zeleny's Method.

Many determinations of the ionic mobilities  $k_1$  and  $k_2$  have been made in various gases at different pressures and temperatures. It is only possible to describe a few of them here.

In 1900 Zeleny made some accurate measurements of  $k_1$  and  $k_2$  for the ions produced by X-rays in several gases at atmospheric pressure.

The gas was passed at an uniform rate through a tube TT' of circular cross-section, along the axis of which was a long insulated cylindrical electrode CF (fig. 1). This electrode was divided into two parts by a narrow gap at DE. A narrow beam of X-rays SAB was passed across the tube from a source S so that positive and negative ions were produced in

the gas over the cross-section AB of the tube. The electrode CF was kept at zero potential and the tube TT' charged to a potential  $V$  by means of a battery. As the ions are carried along the tube in the stream of gas, those of one sign move towards

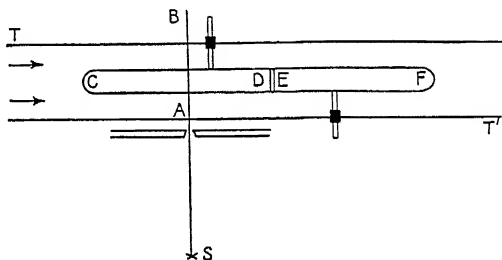


Fig. 1

the tube and the others towards the electrode. The ions produced at the surface of the tube which move to the electrode strike the electrode farthest from AB. If  $V$  is small some ions will reach EF, and as  $V$  is increased the number reaching EF diminishes to zero. The value of  $V$  just big enough to prevent any ions from reaching EF was determined. With this value of  $V$  the ions starting from the surface of the tube at AB move across to E. The ions reaching EF were detected by means of a quadrant electrometer to which it was connected.

Let the radius of the tube be  $b$  and that of the electrode  $a$ , and let  $F$  be the strength of the electric field from the tube towards the electrode. Then  $F$  is inversely as  $r$ , so that  $F = V / (r \log \frac{b}{a})$ , provided there are not enough ions present to sensibly disturb the field. If  $v$  is the velocity of the gas stream, then in a time  $dt$  an ion will be carried along with the stream a distance  $dx = vdt$ , where  $x$  is the distance along the tube from AB, and will move in towards the electrode a distance  $dr = kFdt$ . Hence  $dr = kFdx/v$ , or

$$2\pi v r dr = \frac{2\pi V k dx}{\log(b/a)}.$$

Integrating along the path of the ions from  $r = b$ ,  $x = 0$  to  $r = a$ ,  $x = d$ , where  $d$  is the distance between EF and AB, we get

$$Q = \int_a^b 2\pi v r dr = \frac{2\pi V k}{\log \frac{b}{a}} d,$$

where  $Q$  is the volume of gas flowing past AB in unit time. Hence

$$k = \frac{Q \log b/a}{2\pi V d}.$$

### 3. Langevin's Method.

Another very good method of finding ionic mobilities was used by Langevin. The space between two parallel plates was filled with the gas, and ions were produced by a single flash of X-rays got by breaking the primary circuit of an induction coil connected to the X-ray tube. An electric field was maintained between the plates which was reversed in direction at a time interval  $t$  after the flash. The charge received by one of the plates was measured with a quadrant electrometer. This charge  $Q$  varies with the interval  $t$ , and the ionic mobilities can be found from the relation between  $Q$  and  $t$ . Let the distance between the plates be  $l$  and the strength of the electric field  $X$ . Suppose that the charge on the positive ions produced by the flash of X-rays is  $q$

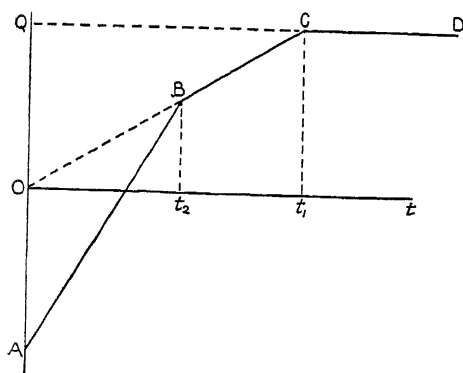


Fig. 2

per unit volume of the gas, and that the positive ions move towards the plate connected to the electrometer during the interval  $t$ . The charge received by the plate during the interval  $t$  will then be  $Aqk_1Xt$ , where  $A$  is the area of cross-section of the flash of X-rays. When the field is reversed the negative ions remaining between the plates will go to the plate, which will therefore receive a negative charge  $Aq(l - k_2Xt)$ , so that we have

$$Q = Aqk_1Xt - Aq(l - k_2Xt) = Aq\{Xt(k_1 + k_2) - l\}.$$

This expression is correct so long as  $k_1Xt$  and  $k_2Xt$  are less than  $l$ . If  $k_2$  is greater than  $k_1$ , as is usually the case, then when  $t$  is greater than  $t_2 = l/k_2X$  we shall have

$$Q = Aqk_1Xt,$$

and when  $t$  is greater than  $t_1 = l/k_1X$  we shall have

$$Q = Aql.$$

The relation between  $Q$  and  $t$  is shown in fig. 2. When  $t = 0$ ,  $Q = -Aql$ , since all the negative ions go to the plate, and when  $t > t_1$  then  $Q = Aql$ , since all the positive ions go to the plate. By measuring  $Q$  for a series of values of  $t$  and plotting the results the values of  $t_1$  and  $t_2$  can be found.  $k_1$  and  $k_2$  are then given by  $k_1 = l/Xt_1$  and  $k_2 = l/Xt_2$ . We have assumed that the charges in the gas are not sufficient to appreciably modify the electric field and that the time  $t$  is so short that no appreciable recombination of the ions takes place. Actually, some recombination does occur, which alters the relation between  $Q$  and  $t$  but does not affect the times  $t_1$  and  $t_2$  corresponding to the breaks in the curve at C and B.

#### 4. Rutherford's Method.

Another excellent method of measuring ionic mobilities was first used by Rutherford.

Two parallel plates at a distance  $l$  apart are used with the gas between them. Ions, of one sign only, are produced at the surface of one of the plates. If now an electric field  $X$  is maintained between the plates for a time  $t$  and then reversed, no ions will reach the other plate unless  $t$  is greater than  $l/kX$ . Thus  $k$  can be determined by finding the value of  $t$  at which the other plate begins to receive a charge. This can be done by varying either  $t$  or  $l$ . The electric field may be applied for the time  $t$  a great many times, the field being reversed, after each application, for long enough to return all ions in the gas to the plate from which they started. In this way the charge to be detected can be greatly increased. The field is applied and reversed at regular intervals by a rotating commutator.

An alternating electric field may be used in this method. Thus if  $X = X_0 \sin(2\pi t/T)$  the distance the ions move is given by

$$\int_0^{T/2} kX dt = kX_0 T/\pi.$$

Hence if  $kX_0 T/\pi = l$  the plate will just begin to receive charge. By varying  $l$  it is easy to find the value of  $kX_0 T/\pi$ .

An error may arise owing to diffusion of the ions. An ion in a gas when there is no electric field does not remain at rest but moves about in an irregular manner first in one direction and then in another. Thus some ions will move a greater distance than  $kX_0 T/\pi$  and some not so far. Those which move farther may not get back to the plate when the field is reversed and so may get still farther from it the next time they move away, and so on. The method therefore tends to give too high a value of the mobility. Zeleny's and Langevin's methods may also be affected by a similar error due to diffusion. This error may

be reduced by superposing on the alternating field a steady field  $\bar{X}$  in the direction tending to retard the motion of the ions from the plate at which they start. In this case

$$X = X_0 \sin \frac{2\pi t}{T} - \bar{X},$$

so that the distance the ions move across is

$$k \int_{t_1}^{t_2} \left( X_0 \sin \frac{2\pi t}{T} - \bar{X} \right) dt.$$

Here  $t_1$  and  $t_2$  are the times between which  $X$  is positive. This is approximately

$$kT \left( \frac{X_0}{\pi} - \frac{\bar{X}}{2} + \frac{\bar{X}^2}{2\pi X_0} \right),$$

when  $\bar{X}$  is not a large fraction of  $X_0$ .

The supply of ions at the surface of one of the plates may be produced by allowing ultra-violet light to fall on the plate. This causes the emission of negative ions by the plate. Another way is to have a hole in the plate covered with fine wire gauze. The gas behind the gauze is ionized by X-rays, and some of the ions of one sign are made to go through the gauze by means of a weak electric field.

## 5. Results of Various Experimenters.

The following table contains some of the results on ionic mobilities which have been obtained.  $k_1$  and  $k_2$  are expressed in centimetres per second for 1 volt per centimetre.

Gas.	Pressure in Mm. of Mercury.	$k_1$ .	$k_2$ .	Method.	Observer.
Air .. ..	760	1.36	1.87	Zeleny's	Zeleny
Hydrogen ..	760	6.70	7.95	"	"
Air .. ..	75	14.8	21.90	Langevin's	Langevin
" .. ..	200	5.45	7.35	"	"
" .. ..	415	2.61	3.31	"	"
" .. ..	760	1.40	1.70	"	"
" .. ..	1430	0.75	0.90	"	"
Carbon monoxide	760	1.10	1.14	"	Wellisch
Carbon dioxide..	760	0.81	0.85	"	"
Nitrous oxide ..	760	0.82	0.90	"	"
Sulphur dioxide	760	0.44	0.41	"	"
Methyl iodide ..	760	0.21	0.22	"	"
Ethyl iodide ..	760	0.17	0.16	"	"
Helium .. ..	760	5.09	6.31	Rutherford's	Franck & Pohl
Argon .. ..	760	1.37	1.70	"	Franck
" (pure) .. ..	760	1.37	206.0	"	"
Nitrogen .. ..	760	1.27	1.84	"	"
" (pure) .. ..	760	1.27	144.0	"	"

It appears that the mobility of the negative ions is generally rather greater than that of the positive ions. The velocity of the positive ions is nearly inversely as the gas pressure. In very pure argon and nitrogen Franck found that the negative ions had much larger velocities than in these gases not specially purified. A small quantity of oxygen added to the pure argon reduced  $k_2$  to a much smaller value.

Zeleny found that water vapour diminishes the velocity of the negative ions in air and hydrogen, making it nearly equal to that of the positive ions. The velocities of the negative ions in air,  $\text{CO}_2$ , and hydrogen at pressures down to a few millimetres have been determined by Lattey and Tizard, using a modification of Rutherford's method. They found that the velocity is approximately a function of  $X/p$  or the ratio of the electric field strength to the pressure. In the case of the positive ions the velocity  $v_1$  is given by  $v_1 = k_1 X/p$ , where  $k_1$  is the mobility at unit pressure. For the negative ions  $v_2 = k_2 X/p$ , when  $X/p$  is small, but as  $X/p$  increases the velocity  $v_2$  increases more rapidly than  $X/p$ .

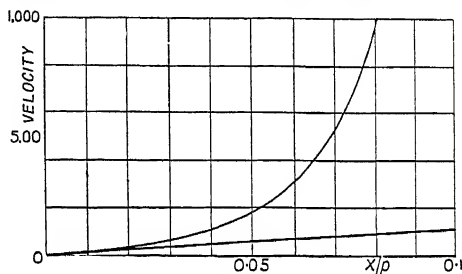


Fig. 3

The relations between  $v_1$  and  $v_2$  and  $X/p$  in dry air are shown in fig. 3. When  $X$  is expressed in volts per centimetre and  $p$  in millimetres of mercury,  $v_1 = 1080X/p$ , and when  $X/p$  is less than about 0.01,  $v_2 = 1350X/p$ , but when  $X/p$  is equal to 0.08 the velocity  $v_2$  is equal to 1000 cm./sec., which is ten times  $1350X/p$ .

## 6. Theory of Ionic Velocities.

It will be convenient now to consider the theory of the motion of ions through a gas in an electric field. The ions are supposed to move about in the gas with a high velocity and to collide with the gas molecules just as the uncharged molecules do. The average kinetic energy of an ion in the absence of an electric field will be equal to the average kinetic energy of a gas molecule, so that we have  $\frac{1}{2}mV^2 = \frac{1}{2}m'V'^2$ , where  $m$  and  $m'$  are the masses of an ion and a molecule, and  $V^2$  and  $V'^2$  the average values of the squares of their velocities.

In an electric field the ions acquire an average velocity of drift  $u$  in the direction of the field. The average momentum of an ion is then  $mu$ , while when there is no field their average momentum is zero, since there are then as many moving in any direction as in the opposite direction. If  $e$  is the charge on an ion then in a field of strength  $X$  the force on it is  $Xe$ , so that the field gives the ion momentum at the rate  $Xe$  per unit time. The ions lose the momentum they receive from the field by colliding with the molecules, and in a steady field the rate at which they lose momentum is equal to that at which they receive it. If  $\lambda$  denotes the mean free path of an ion, or the average distance between its collisions with the gas molecules, then the number of collisions it makes in unit time is approximately  $V/\lambda$ . This is not exact, because  $V$  is not the average velocity but the square root of the mean square of the velocity. If on the average an ion loses a fraction  $f$  of its average momentum  $mu$  at each collision, the momentum it loses in unit time is  $Vfmu/\lambda$ . Hence in a steady state

$$Xe = Vfmu/\lambda,$$

so that

$$u = \frac{Xe\lambda}{fmV}.$$

In the case of an ion having a mass very small compared with that of a gas molecule,  $f$  will be nearly unity, so that

$$u = \frac{Xe\lambda}{mV},$$

but in the case of a heavy ion having a mass larger than that of a gas molecule  $f$  will be a small fraction.

## 7. Mobility and Coefficient of Diffusion.

The mobility of an ion may be obtained from its coefficient of diffusion  $K$ . We have

$$-K \frac{dp}{dx} = pv,$$

where  $p$  denotes the partial pressure of the ions, and  $v$  the velocity of diffusion in the  $x$  direction. Also if there are  $n$  ions per unit volume —  $dp/dx$  may be regarded as the force on the ions in unit volume which drives them along with the velocity  $v$ . In an electric field  $X$  the force on the  $n$  ions is  $Xen$ , and this gives them a velocity  $kX$ . Hence, assuming the velocity proportional to the driving force, we have

$$\frac{v}{kX} = \frac{-dp/dx}{Xen} = \frac{pv}{KXen},$$

or

$$k = \frac{Ken}{p}.$$

But  $p = nR_1T$ , where  $R_1$  is the gas constant for one molecule, so that

$$k = \frac{Ke}{R_1T}.$$

If  $\mathcal{N}$  is the number of molecules in 1 gm. molecule or mol, then

$$k = \frac{K\mathcal{N}e}{RT},$$

where  $R$  is the gas constant for 1 mol. We have  $\mathcal{N}e = 9650$  electromagnetic units, and  $R = 8.32 \times 10^7$ , so that at  $300^\circ \text{K}$ . we get, after multiplying by  $10^8$  to convert electromagnetic units of field strength into volts per centimetre,  $k = 39K$ . The coefficient of diffusion of oxygen molecules diffusing in oxygen at 760 mm. and  $300^\circ \text{K}$ . is approximately 0.19, which gives  $k = 7.4$ . For hydrogen diffusing through hydrogen  $K = 1.3$ , so that  $k = 51$ . The values found for  $k_1$  in oxygen and hydrogen are 1.4 and 6.7, and for  $k_2$  1.9 and 8.0 cm. per second for 1 volt per centimetre. These values are from four to eight times smaller than the values just calculated. This is believed to indicate that the ions are not single molecules carrying a charge  $e$  but clusters of several molecules held together by the electrical field of the charge.

In the case of the negative ions the clusters evidently diminish in size as  $X/p$  increases, and when  $X/p$  is greater than about 0.1 the negative ions are reduced to electrons. The motion of electrons in gases is considered in another chapter.

## 8. Townsend's Determination of Coefficients of Diffusion.

The coefficients of diffusion of the ions produced in gases by X-rays or other sources of ionization were determined by J. S. Townsend. Townsend passed the gas containing ions through narrow metal tubes, and determined the fraction

of the ions lost by diffusion to the walls of the tubes. He also determined the ratio of the mobility to the coefficient of diffusion by the method described in the chapter on the motion of electrons in gases. This ratio, as we have seen above, is equal to  $\mathcal{N}e/RT$ . If  $e$  is equal to the charge on one univalent ion in a solution then we know that  $\mathcal{N}e$  is equal to 96,500 coulombs, so that Townsend's determinations of  $k/K$  enabled him to prove that the charges on gaseous ions are equal to the charge on a univalent ion in solutions. He found  $\mathcal{N}e$  for positive ions always nearly equal to 96,500 coulombs, and the same was true for negative ions when they had small mobilities, showing that they were clusters of molecules like the positive ions. The values of the coefficients of diffusion of ions found by Townsend by the narrow tube method are as follows for ions produced by X-rays:

Gas.	Positive Ions.	Negative Ions.
Air .. ..	0.028	0.043
Oxygen .. ..	0.025	0.040
Carbonic acid ..	0.023	0.026
Hydrogen ..	0.123	0.190

Townsend also measured the coefficient of diffusion of the ions produced in air by several different sources of ionization and found the following values:

	Positive Ions.	Negative Ions.
X-rays .. ..	0.028	0.043
Radium rays ..	0.032	0.043
Ultra-violet light ..	—	0.043
Point discharge ..	0.024	0.035

He found that the coefficients of diffusion were inversely as the gas pressure from 200 mm. to 772 mm. of mercury.

Salles has made measurements of the coefficients of diffusion of ions in several gases by Townsend's tube method, and obtained results agreeing approximately with those found by Townsend.

### 9. Another Method of Determining the Charge on One Mol of Gaseous Ions.

The charge carried by 1 gm.-molecule or mol of gaseous ions was determined by the writer by an entirely different method. Air containing a small amount of a solution of an alkali salt in suspension in the form of fine spray was passed through a long platinum tube heated in a furnace to a high temperature. The salt is volatilized in the tube and is ionized at the high temperature. The charge carried by the ions formed was found by measuring the current between the platinum tube and a cylindrical electrode placed along the axis of the tube. It was found that above about 1300° C. and with a potential difference between the electrode and tube of more than 600 volts the current obtained was nearly independent of the temperature and potential difference. This saturation current was measured for a number of different alkali salts, and was found to be proportional to the amount of salt passing into the tube in unit time and inversely

proportional to the electrochemical equivalent of the salt. The quantity of electricity carried by the ions formed from 1 gm. equivalent of any of the salts used was found to be approximately 98,000 coulombs, which agrees with the value 96,500 found in solutions within the limits of error. The salts used were  $\text{CsCl}$ ,  $\text{Cs}_2\text{CO}_3$ ,  $\text{RbI}$ ,  $\text{Rb}_2\text{CO}_3$ ,  $\text{RbCl}$ ,  $\text{KCl}$ ,  $\text{KI}$ ,  $\text{KBr}$ ,  $\text{KF}$ ,  $\text{K}_2\text{CO}_3$ ,  $\text{NaI}$ ,  $\text{NaCl}$ ,  $\text{NaBr}$ ,  $\text{Na}_2\text{CO}_3$ ,  $\text{LiI}$ ,  $\text{LiCl}$ ,  $\text{LiBr}$ ,  $\text{Li}_2\text{CO}_3$ .

The measurements which have been made on the ratio of the charge to the mass of positive rays, which are discussed in the chapter on positive rays, show that the positive ions formed in gases at low pressures may have charges which are small multiples of the ionic charge  $e$ , but in gases at atmospheric pressure at any temperature gaseous ions are found to have charges equal to  $e$ . Probably in electric discharges with very intense fields ions having charges which are small multiples of  $e$  may be formed even at high pressures.

The fact that ions formed under such varied conditions in liquid electrolytes of any kind, in gases by any source of ionization, at any pressure and at any temperature and from any kind of compound or element, always have charges which are small but exact multiples of the ionic charge  $e$ , shows conclusively that electricity has an atomic constitution. That is to say that electricity is composed of small exactly equal atoms of electricity and that the charge on the negative atoms is equal to the charge on the positive atoms.

## 10. Recombination of Ions.

When positive and negative ions are present in a gas they recombine and form electrically neutral molecules. The rate of recombination is proportional to the number  $n_1$  of positive ions and to the number  $n_2$  of negative present in unit volume. Thus we have

$$\frac{dn_1}{dt} = \frac{dn_2}{dt} = -\alpha n_1 n_2,$$

where  $\alpha$  is a constant called the coefficient of recombination. If  $n_1 = n_2 = n$ , then

$$\frac{dn}{dt} = -\alpha n^2,$$

which gives  $\frac{1}{n} - \frac{1}{n_0} = \alpha t$ , where  $n_0$  is the value of  $n$  when  $t = 0$ . We are supposing here that no ions are removed from the gas except by recombination.

The coefficient of recombination of ions in gases has been determined by Rutherford, Townsend, Langevin, and others by measuring the rate at which the ions disappear. The values of  $\alpha/e$  found in different gases at atmospheric pressure vary between 3000 in hydrogen and 3400 in air, with  $e$  in electrostatic units. The factor  $e^{-1}$  appears because the quantity actually measured is not the number of ions  $n$  but the charge of  $q = ne$  which they carry. Thus we have

$$\frac{e}{q} - \frac{e}{q_0} = \alpha t,$$

so that

$$\frac{\alpha}{e} = \frac{1}{t} \left( \frac{1}{q} - \frac{1}{q_0} \right).$$

Langevin found that  $\alpha/e$  is nearly proportional to the pressure of the gas between 760 and 150 mm. of mercury. At pressures of several atmospheres in air and  $\text{CO}_2$  he found  $\alpha/e$  to be nearly inversely as the pressure. At very low pressures the coefficient of recombination probably becomes very small, but no determinations of it have been made at low pressures.

A simple theory of recombination has been given by Langevin which is approximately correct in gases at high pressures. Consider a sphere of radius  $r$  in a gas with a positive ion at its centre. If there are  $n_2$  negative ions per unit volume, the number of negative ions entering the sphere per unit time owing to the electric field of the positive ion will be  $4\pi r^2 n_2 (k_1 + k_2) \frac{e}{r^2}$ , since a positive and a negative ion move towards each other with the relative velocity  $(k_1 + k_2) \frac{e}{r^2}$ , and when  $r$  is small there will very seldom be more than one negative ion at the surface of the sphere. If there are  $n_1$  positive ions present in unit volume the total number of negative ions moving up to positive ions per unit volume in unit time will therefore be  $4\pi e n_1 n_2 (k_1 + k_2)$ , so that if we suppose that recombination occurs in every case we get

$$\alpha = 4\pi e (k_1 + k_2).$$

At high pressure this formula gives values of  $\alpha/e$  in agreement with those observed. For example, in carbonic acid Langevin found that at three atmospheres pressure

$$\frac{\alpha}{4\pi e (k_1 + k_2)} = 0.97.$$

At pressures below one atmosphere  $\alpha$  is much smaller than  $4\pi e (k_1 + k_2)$ . This is due to the fact that an ion may move up to another one and move round it and then diffuse away without recombining. Unless the ion loses kinetic energy by collisions when close to the other ion it is unlikely to recombine. The electric field  $e/r^2$  of an ion is too weak, except at very small distances, to appreciably affect the diffusion of another ion.

## 11. Formation of Clouds on Ions.

Clouds consisting of minute drops of water are formed in gases by the condensation of supersaturated water vapour on dust particles or other nuclei. In the absence of such nuclei no cloud is formed unless the degree of supersaturation is very high. C. T. R. Wilson discovered that the ions formed in gases by X-rays or other sources of ionization can act as nuclei for the condensation of supersaturated water vapour. The apparatus used by C. T. R. Wilson is shown in fig. 4. BB is a glass tube about 20 cm. long and 4 cm. in diameter which is closed at its lower end by a rubber stopper R. Another glass tube CC with its upper end closed slides freely up and down inside BB. A tube DD passing through the stopper connects the inside of CC through a valve S, opened by pulling T, to a large bottle E, which is kept exhausted by means of a tube P leading to a pump. The tube BB is about half filled with water as shown.

The upper end of BB is connected to a mercury manometer F through a stopcock, and air can be withdrawn or admitted through a tube G packed with glass wool to remove dust particles. A bulb A is connected to the top of BB by means of a wide tube. If the valve S is shut and the stopcock Q opened, the sliding tube CC, which acts as a piston, may be raised to any desired point by removing some air through G. If Q is then shut and S opened a vacuum is produced under

the piston, which very suddenly moves down until it strikes the rubber stopper. In this way the air in A is made to expand very rapidly. The change of pressure due to the sudden expansion can be measured with the manometer F. The expansion produces supersaturation of the water vapour in A, and if any dust particles are present a cloud is formed which falls slowly and finally settles on the glass walls, so removing the dust particles from the gas.

By repeating this process two or three times all dust can be got rid of. In dust-free air no cloud is formed when the expansion ratio

is less than 1.25. With ratios between 1.25 and 1.38 a small number of drops, about 100 per cubic centimetre, appear, and with expansion ratios above 1.38 a very dense cloud is formed which is evidently due to condensation on the molecules.

If X-rays are passed through the bulb A then a cloud is formed with expansion ratios between 1.25 and 1.38. This effect persists for a short time after the rays are cut off. If an electrode is sealed into A, it is found that charging it to a few hundred volts stops the formation of any cloud after the rays are cut off.

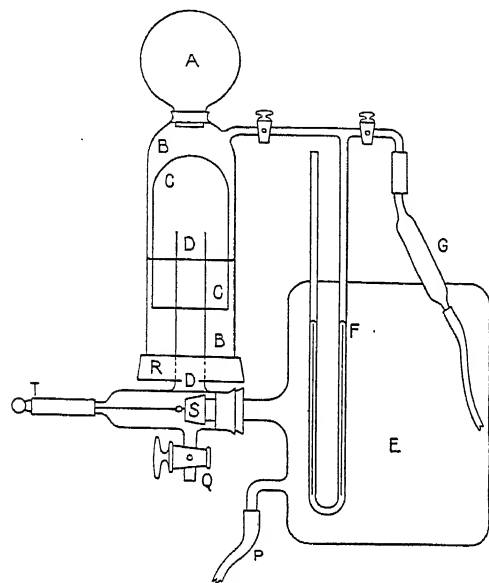


Fig. 4

This shows that the cloud produced by the rays is due to condensation on the ions formed by the rays. C. T. R. Wilson found that the clouds are formed more easily on negative than on positive ions. An expansion of 1.25 gives a cloud with negative ions, but 1.34 is required with positive ions.

## 12. Photographs of Tracks of $\alpha$ - and $\beta$ -rays.

C. T. R. Wilson found that it is possible to make visible and to photograph the tracks of individual  $\alpha$ -rays and  $\beta$ -rays through gases by forming a cloud on the ions produced by the rays. For this purpose a special form of the expansion apparatus is used. The expansion chamber is a tube about 15 cm. in diameter closed at the top with a glass plate. The top of the piston is also a flat plate, painted black

and nearly as large as the top of the expansion chamber. The distance between the top of the expansion chamber and the top of the piston is about 3 cm., and is suddenly increased when the piston is connected to the vacuum bottle as with the first apparatus. A rather large potential difference is maintained between the top and bottom of the expansion chamber to remove ions formed immediately. If a very minute particle of radium is supported on a rod inside the expansion chamber, then when the sudden expansion occurs the tracks of the  $\alpha$ -rays appear as long narrow lines of cloud. To see them clearly it is necessary to illuminate the chamber strongly. The tracks of the  $\alpha$ -rays are usually straight lines, but occasionally a track appears in which there is a sudden change of direction at one point which is usually near the end of the track. These tracks must be photographed immediately after they have been formed. They very quickly disappear. Two photographs are reproduced here of  $\alpha$ -ray tracks through nitrogen, showing collision between an  $\alpha$ -ray and a nitrogen atom. (See fig. 5, Plate facing p. 131.)

Harkins and also Chadwick have taken photographs of enormous numbers of  $\alpha$ -ray tracks. Chadwick obtained several photographs showing collisions between an  $\alpha$ -ray and a nitrogen atom in which a hydrogen atom was knocked out of the nitrogen atom. The hydrogen atoms gave straight tracks, much narrower than the  $\alpha$ -ray tracks and easily distinguishable from them. In such collisions the  $\alpha$ -ray apparently remains inside the nitrogen atom and the nitrogen atom acquires sufficient velocity to produce a short track. The track of the  $\alpha$ -ray therefore branches at the collision into a long straight very narrow track and a short thick track. Collisions are frequently observed in which an  $\alpha$ -ray track branches into two similar short tracks. This is supposed to be due to the  $\alpha$ -ray colliding with an atom and rebounding from it, so that the  $\alpha$ -ray and the atom both produce a short track after the collision. By measuring the lengths of the branches the velocities of the colliding particles can be estimated, and it is found that in these collisions energy and momentum are conserved as for perfectly elastic bodies. When a narrow beam of X-rays is examined by this method two distinct types of track are observed. The electrons shot out of the atoms give tracks about 1 cm. or so long which are not straight but continually change in direction, especially towards the end of the track. These tracks start out sideways from the X-ray beam and more often with a forward velocity component than not. Besides these rather long tracks very short tracks are also observed which are believed to be due to the scattering of an X-ray quantum by an electron. The electron then gets only a small amount of energy and so only gives a short track. A short track is frequently associated with a long track which starts outside the X-ray beam and is supposed to be produced by the scattered quantum.

Thus C. T. R. Wilson's method enables effects due to single atoms and electrons to be observed and photographed, and has already led to most interesting and important results.

### *Direct Determination of the Ionic Charge*

13. Up to about 1897 the value of the ionic charge  $e$  could only be obtained from the number of molecules in gases, as deduced from the viscosity and other properties of gases by the results of the kinetic theory. The product  $\mathcal{N}e$ , where  $\mathcal{N}$  is the number of molecules in 1 gm.-molecule or mol of any gas, was known accurately from the measurements of the electrochemical equivalent of silver, but the possible error in the value of  $\mathcal{N}$  deduced from the kinetic theory of gases was 50 per cent or more. About that date there was nevertheless convincing evidence available proving the atomic nature of electricity. The facts of electrolysis alone showed that the average ionic charge in liquid electrolytes is always a small integral multiple of a definite unit of charge, and the fact that the rays in a beam of cathode rays are all deflected to the same extent by a magnetic field showed that the rays all have the same value of  $e/m$  and are not a mixture of different sorts of particles having constant properties only on the average. About that time it was shown by J. J. Thomson and others that electrons obtained from many different sources have identical properties, and it was clear that these electrons are constituents of the chemical atoms, which were therefore not indivisible particles. These conclusions were supported by an immense amount of new evidence during the next ten years, and the theory of the atomic nature of electricity was established beyond reasonable doubt before Millikan succeeded in measuring very small electric charges with sufficient accuracy to show that they were always exact multiples of an atomic unit. This result came therefore as a final confirmation of a well established theory, but was and is nevertheless of the highest interest and importance.

### **14. Townsend's Method.**

The first experiments which could be regarded as a direct determination of the ionic charge were made by Townsend in 1897. The results he obtained were not very exact, but the method he devised contained several of the essential features of the methods used in subsequent investigations. Townsend showed that it was possible to make a direct determination of the ionic charge, and started the attack on the problem.

Townsend found that the gases evolved in the electrolysis of dilute sulphuric acid by rather large currents form dense clouds when bubbled through water. These clouds consist of minute drops of water containing a trace of some hygroscopic substance, probably  $\text{H}_2\text{SO}_4$ , and they are electrically charged. He measured the charge carried by

the cloud and the amount of water in it, and found the mass of the droplets by measuring the rate at which they fell through the gas. In this way he got the total number of drops in the cloud and the total charge on them. Dividing the charge by the number of drops he got the average charge per drop, which was about  $3 \times 10^{-10}$  electrostatic units. Later he found that all the drops were not charged with electricity of the same sign, and allowing for this got for the average charge per drop  $5 \times 10^{-10}$  e.s.u. Townsend supposed that the drops were formed on gaseous ions, and so regarded the result obtained as a determination of the ionic charge. The drops in Townsend's clouds all fell at nearly the same rate, since the upper surface of the cloud remained distinct as it fell. This showed that the majority of the drops were of equal size, and so justified to some extent the assumption that they all carried equal charges.

The size of the drops was obtained from the rate of fall by means of the theory, due to Stokes, of the motion of a sphere through a viscous fluid. According to Stokes's theory the resistance to the motion of a sphere of radius  $a$  through a medium of viscosity  $\mu$ , with small uniform velocity  $v$ , is equal to  $6\pi\mu va$ . It is assumed in deducing this expression that there is no slipping at the surface of the sphere and that  $av\rho'/\mu$  is a small fraction,  $\rho'$  being the density of the medium. For a sphere of water falling through air,  $a$  must be small compared with 0.006 cm. The weight of a sphere of density  $\rho$  is  $\frac{4}{3}\pi a^3 \rho g$ , so that for a falling sphere we have

$$\frac{4}{3}\pi a^3 g(\rho - \rho') = 6\pi\mu va,$$

when the velocity  $v$  has attained a constant value. Hence

$$v = \frac{2}{9} \frac{\rho - \rho'}{\mu} ga^2,$$

which is the expression used by Townsend to get the radius  $a$  and so the mass  $m = \frac{4}{3}\pi a^3 \rho$  of his droplets.

The weak points in Townsend's determination of  $e$  are the assumption that all the drops carry equal charges and that these equal charges are equal to the ionic charge. It was not clear that only one ion went into each drop—in fact the way in which the charges got into the drops was not known.

### 15. Method of J. J. Thomson.

Soon after Townsend's experiments appeared, J. J. Thomson carried out a determination of  $e$  by a method which was based on essentially the same principles, but in which the droplets were formed by condensing water vapour on the ions formed in air by X-rays or radium rays.

It had been shown by C. T. R. Wilson that such ions act as nuclei for the condensation of [water vapour, so that in J. J. Thomson's experiments it was clear that the charge measured was the ionic charge.

This celebrated investigation has therefore sometimes been regarded as the first direct determination of the ionic charge. J. J. Thomson obtained the cloud on the ions by means of C. T. R. Wilson's expansion apparatus described above, and got the mass of the drops from their rate of fall. He found the charge in the gas by measuring its electrical conductivity due to a small electric field. The conductivity is equal to  $ne(k_1 + k_2)$ , where  $n$  is the number of positive or negative ions per cubic centimetre, and  $k_1$  and  $k_2$  are the mobilities of the ions. The mobilities in air were known and so  $ne$  could be calculated. The total mass of water in the cloud was calculated from the expansion ratio, assuming the expansion to be adiabatic. In this way J. J. Thomson showed that  $e$  was nearly the same for ions produced by X-rays, radium rays, and ultra-violet light. The final result he got for  $e$  was  $3.4 \times 10^{-10}$  electrostatic units.

The weak points in these experiments are the assumptions that all the drops contain only one ion and that the total mass of water in the cloud is equal to that due to an adiabatic expansion. As a matter of fact the clouds soon evaporate as the air warms up, so that the mass of the drops is not constant.

#### 16. Another Method of Determining the Ionic Charge.

A method of finding the charge on the drops which makes it unnecessary to determine the total mass of the cloud and the total number of drops was proposed by the writer in 1903. The velocity of a sphere moving through a viscous liquid is proportional to the force driving it, so that in a vertical electric field  $F$  which exerts a force on the drops equal to  $Fe$  the velocity of the drops will be changed. If  $v_1$  is the rate of fall with  $F = 0$ , and  $v_2$  the rate of fall in the field, then

$$\frac{v_1}{v_2} = \frac{mg}{mg + Fe},$$

where  $m$  is the mass of the drop, and the force  $Fe$  is reckoned positive when its direction is downwards. This gives

$$e = \frac{mg(v_2 - v_1)}{Fv_1}.$$

The mass  $m$  can be got from  $v_1$  as in Townsend's and J. J. Thomson's experiments, so that  $e$  can be calculated from  $v_1$  and  $v_2$ .

The apparatus used consisted of a C. T. R. Wilson expansion chamber containing two horizontal parallel electrodes between which a vertical electric field could be maintained by means of a battery giving up to 2000 volts. The electrodes were 3.5 cm. in diameter and about 5 mm. apart. The space between them was illuminated by a narrow parallel beam, and the cloud was observed on the axis of the electrodes where the field was practically uniform. The moist air in the expansion

chamber was ionized by X-rays, which were usually cut off just before an expansion.

It was found that the clouds formed with no electric field consisted of drops which very nearly all fell at practically the same rate. The individual drops could be seen unless the cloud was very dense, and it was easy to see if there were any drops falling at different rates. This was not the case in the cloud near the axis where it was observed.

It was also found that the clouds formed in successive expansions under the same conditions all fell at practically the same rate, so that the plan adopted was to measure the time for the top of a cloud to fall from the upper electrode to the lower one, first without any field and then with a field. This gave better results than were obtained by measuring the rate of fall for part of the distance without any field and for the rest of the distance with a field applied. Owing to the evaporation of the drops the rate of fall is not uniform, so that it was better to measure the average rate over the whole distance without any field and then with a field. The fact that the clouds formed in successive expansions all fell at the same rate justified this procedure, which was adopted because it gave the best results and not because the method of measuring  $v_1$  and  $v_2$  on the same cloud was not tried.

It was found that in an electric field all the drops did not fall at the same rate. Several sets of drops could be seen and all the drops in each set fell at the same rate. Three such sets could usually be detected, and it was found that the charges on the drops in them were nearly in the ratio 1:2:3. This result showed clearly that the charges on the drops were multiples of an atomic unit, as they should be according to the atomic theory of electricity.

If the X-rays were kept on during and after the expansion a few droplets were obtained carrying comparatively large charges. Some of these drops could be made to rise with a potential difference of only a few hundred volts. The charges on these drops were found to be rather large multiples of the ionic charge and they were not used to determine  $e$ , because owing to evaporation it did not seem to be possible to determine their charges with sufficient accuracy to establish the value of the integer expressing the charge in terms of  $e$ . It was only possible to be sure of the correct value of the integer when it was not greater than two or three. Owing principally to the evaporation of the drops it was not possible to obtain accurate results. The results obtained varied from  $2 \times 10^{-10}$  to  $4.4 \times 10^{-10}$  electrostatic units. The mean result was  $3.1 \times 10^{-10}$ .

### 17. Millikan's Method.

The problem of making an accurate direct determination of the ionic charge was finally solved by Millikan about 1908. His method was in principle the same as that used by the writer, but he eliminated

the error due to evaporation by using small drops of oil or mercury, which do not evaporate, and by using a strong electric field and drops carrying several ionic charges he was able to make the drops move up or down and so keep a single drop under observation for a long time and make a series of measurements with it. Millikan made a long series of measurements, taking all possible precautions to eliminate errors, and his final result is believed to be correct to one part in one thousand.

Millikan's final form of apparatus is shown diagrammatically in fig. 6. AA and BB are two circular metal plates with optically worked plane surfaces. These plates are separated by three glass blocks cut from a piece of plane parallel optically worked glass. The distance

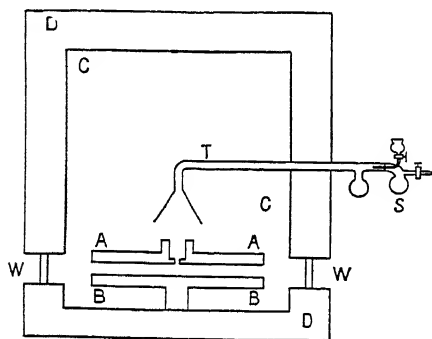


Fig. 6

between the plates is 14.9174 mm. At the centre of the upper plate there is a very small hole as shown. The tube T leads to a sprayer S by means of which oil or mercury spray can be produced and blown into the space above the plates. The plates are supported inside a metal box CC immersed in an oil tank DD. Glass windows WW enable a beam of light to be passed between the plates so as to illuminate the

drops at the centre of the plates. The drops were observed through a telescope in a direction slightly inclined to the beam of light. X-rays could also be passed between the plates when desired. The plates could be connected to a battery giving up to 10,000 volts.

On working the sprayer for a few seconds a large number of drops is formed above the plates, and a few of these fall through the small hole into the space between the plates. The drops are charged, and usually a few have such masses and charges that they remain suspended in the electric field or move very slowly up or down. A drop which moves slowly up may be selected, and its small upward velocity is determined by measuring with a chronograph the time the image of the drop seen in the telescope takes to go from one cross hair in the eyepiece to another. The images of the drops are seen as minute points of light like stars, and the instant at which they pass a cross hair can be very accurately timed. The electric field is then cut off and the velocity with which the drop falls is determined. The field is then put on and the upward velocity again found. In this way a single drop can be kept under observation for several hours and a

great many observations made on it. The charge on the drop can be varied when desired by passing X-rays between the plates. The air is ionized by the rays and an ion usually gets on to the drop very soon. The change in the charge on the drop when an ion gets on it is immediately made evident by the change in the velocity of the drop in the electric field.

For a particular drop the charge on it is proportional to  $v_2 - v_1$ , the difference between its velocity in the field and that due to gravity alone. When  $v_2$  is an upward velocity the charge is proportional to the sum of the numerical values of the two velocities. The following table gives a set of numbers equal to  $v_2 - v_1$  multiplied by a constant ( $A$ ) which were obtained by Millikan in a series of observations on one oil drop. The charge on this drop was changed from time to time and its velocities were found.

$A(v_2 - v_1)$		$A(v_2 - v_1)$
$19.66 \div 4 = 4.915$		$53.91 \div 11 = 4.901$
$24.60 \div 5 = 4.920$		$59.12 \div 12 = 4.927$
$29.62 \div 6 = 4.937$		$63.68 \div 13 = 4.898$
$34.47 \div 7 = 4.924$		$68.65 \div 14 = 4.904$
$39.38 \div 8 = 4.922$		$78.34 \div 16 = 4.896$
$44.42 \div 9 = 4.935$		$83.22 \div 17 = 4.895$
$49.47 \div 10 = 4.947$		

All the numbers found for  $A(v_2 - v_1)$  when divided by an integer give nearly equal numbers. This shows that the charges on the drop were always exact multiples of a definite unit, which was the ionic charge, because the charge was varied by adding positive or negative ions one at a time. These results show that all the ions carry the same charge either positive or negative, so that the ionic charge is not an average value for a quantity which varies, but a definite atomic unit.

The absolute value of  $e$  was found from a very large number of measurements on many different drops. The viscosity of dry air was very carefully redetermined, and also of course the exact density of the oil used.

It was found that Stokes's law for a falling drop,

$$v = \frac{2}{9} \frac{\rho - \rho'}{\mu} g a^2,$$

is not exactly true for very small drops, and the deviation was accurately determined and allowed for. The final result obtained was

$$e = 4.774 \times 10^{-10} \text{ electrostatic units.}$$

The ionic charge has been determined by several other methods in recent years. Rutherford and Geiger determined the charge  $2e$  on  $\alpha$ -rays and got  $e = 4.65 \times 10^{-10}$  and Regener by the same method

got  $4.79 \times 10^{-10}$ . This method is described in the chapter on Cathode Rays,  $\beta$ -rays, and  $\alpha$ -rays.

### 18. Perrin's Investigations. Brownian Movements and Diffusion.

Perrin has made a very interesting series of investigations on the Brownian movements of small particles suspended in liquids, from which he has deduced the number of molecules in unit volume of any gas and hence the value of the ionic charge. Very small particles suspended in a liquid continually move about in an irregular manner visible in a high-power microscope. This motion is believed to be due to collisions between the particles and the molecules of the liquid. The particle may be regarded as a big molecule, and its average kinetic energy should be equal to the average kinetic energy of a gas molecule at the same temperature.

An emulsion containing  $n$  equal particles per unit volume suspended in a liquid may be compared with a gas containing  $n$  molecules in unit volume. The particles may be regarded as exerting a pressure equal to  $\frac{1}{3}mnV^2$ , where  $m$  is the mass of a particle, and  $V^2$  is the average value of the square of the velocities of the particles. If we consider a horizontal layer of thickness  $dx$  in the emulsion, the downward force of gravity on the  $ndx$  particles in unit area of the layer must be balanced by the upward force arising from the variation of the gas pressure  $p$  due to the motion of the particles. If  $\rho'$  is the density of the liquid and  $\rho$  that of the particles, we have therefore

$$\frac{nmg(\rho - \rho')}{\rho} dx = -\frac{\partial p}{\partial x} dx.$$

Putting  $p = \frac{1}{3}mnV^2$ , we get

$$\frac{1}{n} \frac{\partial n}{\partial x} = -3g \frac{\rho - \rho'}{\rho V^2}.$$

Integrating this, and putting  $n = n_0$  at  $x = 0$ , we find

$$\log \frac{n_0}{n} = 3g \frac{\rho - \rho'}{\rho V^2} x.$$

Perrin prepared an emulsion of equal gamboge particles and placed a small quantity of it in a shallow glass cell. The particles in it were observed with a vertical microscope focused on a horizontal plane in the emulsion. The number of particles visible in the focal plane was counted, and by moving the microscope up and down through known distances the variation of this number with the level of the focal plane was determined. This gave  $\frac{1}{x} \log \frac{n_0}{n}$  and so  $V^2$  could be calculated by means of the above equation. The densities  $\rho$  and  $\rho'$  and the size of the particles were also determined, so that the mass  $m$  and hence  $mV^2$

for the particles could be calculated. Now  $mV^2$  is equal to  $3p/n$  where  $p$  is the pressure of any gas containing  $n$  molecules in unit volume. Thus Perrin was able to calculate the number of molecules in unit volume of any gas at 760 mm. pressure and at the temperature of his experiments. From this the number  $\mathcal{N}$  of molecules in a gramme-molecule or mol immediately follows, and since  $\mathcal{N}e = 9650$  the ionic charge  $e$  can be deduced. In this way Perrin got  $e = 4.2 \times 10^{-10}$  electrostatic units.

Perrin also observed the diffusion of the particles through the liquid. If  $n$  is the number of particles in unit volume and  $K$  the coefficient of diffusion we have

$$-K \frac{\partial n}{\partial x} = un,$$

where  $u$  is the velocity of diffusion in the  $x$  direction. If  $v$  and  $w$  are the velocity components due to diffusion in the  $y$  and  $z$  directions we have also

$$-K \frac{\partial n}{\partial y} = vn,$$

$$-K \frac{\partial n}{\partial z} = wn.$$

But 
$$\frac{\partial}{\partial x}(un) + \frac{\partial}{\partial y}(vn) + \frac{\partial}{\partial z}(wn) = -\frac{\partial n}{\partial t},$$

so that 
$$K \left( \frac{\partial^2 n}{\partial x^2} + \frac{\partial^2 n}{\partial y^2} + \frac{\partial^2 n}{\partial z^2} \right) = \frac{\partial n}{\partial t}.$$

Now consider a large number  $N$  of particles distributed in any manner over a certain region in a liquid, and let them all be very far from the boundary of the liquid. The particles will diffuse about so that the region they occupy will gradually get larger. Consider the mean value of the square of the distance  $R$  of a particle from any fixed point, and take this point as the origin, so that  $R^2 = x^2 + y^2 + z^2$ . The mean value of  $R^2$  is given by

$$\bar{R}^2 = \frac{\int (x^2 + y^2 + z^2) n \, dx \, dy \, dz}{\int n \, dx \, dy \, dz},$$

where the integration is extended over all the space where  $n$  is not zero. Differentiating with respect to the time  $t$  we get

$$N \frac{\partial \bar{R}^2}{\partial t} = \int (x^2 + y^2 + z^2) \frac{\partial n}{\partial t} \, dx \, dy \, dz,$$

since  $N = \int n \, dx \, dy \, dz$ .

Putting  $K \left( \frac{\partial^2 n}{\partial x^2} + \frac{\partial^2 n}{\partial y^2} + \frac{\partial^2 n}{\partial z^2} \right)$  for  $\frac{\partial n}{\partial t}$ , we obtain

$$\begin{aligned} N \frac{d\bar{R}^2}{dt} &= K \int (x^2 + y^2 + z^2) \left( \frac{\partial^2 n}{\partial x^2} + \frac{\partial^2 n}{\partial y^2} + \frac{\partial^2 n}{\partial z^2} \right) dx dy dz \\ &= K \int \left( x^2 \frac{\partial^2 n}{\partial x^2} + y^2 \frac{\partial^2 n}{\partial y^2} + z^2 \frac{\partial^2 n}{\partial z^2} \right) dx dy dz + \\ &K \int \left\{ \frac{\partial}{\partial x} \left( (y^2 + z^2) \frac{\partial n}{\partial x} \right) + \frac{\partial}{\partial y} \left( (x^2 + z^2) \frac{\partial n}{\partial y} \right) + \frac{\partial}{\partial z} \left( (x^2 + y^2) \frac{\partial n}{\partial z} \right) \right\} dx dy dz. \end{aligned}$$

The second integral may be transformed into a surface integral by Green's theorem, and, since it is to be taken over a volume so large that  $n$  is zero near the surface enclosing the volume, this integral is zero.

Integrating  $\int x^2 \frac{\partial^2 n}{\partial x^2} dx$  by parts gives

$$\left[ x^2 \frac{\partial n}{\partial x} \right] - 2 \int \frac{\partial n}{\partial x} x dx = -2 \int x dn,$$

since  $\left[ x^2 \frac{\partial n}{\partial x} \right]$  is zero at both limits.

$$\text{Again} \quad -2 \int x dn = -2 [nx] + 2 \int n dx = 2 \int n dx,$$

since  $[nx]$  is zero at the limits.

$$\text{In the same way} \quad \int y^2 \frac{\partial^2 n}{\partial y^2} dy = 2 \int n dy,$$

$$\text{and} \quad \int z^2 \frac{\partial^2 n}{\partial z^2} dz = 2 \int n dz.$$

$$\text{Hence} \quad K \int \left( x^2 \frac{\partial^2 n}{\partial x^2} + y^2 \frac{\partial^2 n}{\partial y^2} + z^2 \frac{\partial^2 n}{\partial z^2} \right) dx dy dz = 6K \int n dx dy dz,$$

$$\text{so that we get} \quad \frac{d\bar{R}^2}{dt} = 6K.$$

Since  $R^2 = x^2 + y^2 + z^2$ , we see that the mean square of the distances of the particles from a straight line will increase with the time at the rate  $4K$ , and the mean square of the distances from a plane at the rate  $2K$ . The equation defining the coefficient of diffusion  $K$ ,

$$-K \frac{\partial n}{\partial x} = nu,$$

gives, since the gas pressure  $p$  due to the particles is proportional to  $n$ ,

$$-K \frac{\partial p}{\partial x} = pu = \frac{1}{3} m n u V^2.$$

But  $-\partial p/\partial x$  may be regarded as the force on the  $n$  particles in unit volume which causes them to move with the velocity  $u$ , so that if  $s$  denotes the average velocity with which the particles fall through the liquid under the force of gravity  $m \frac{\rho - \rho'}{\rho} g$ , we have

$$-\frac{\partial p}{\partial x} = \frac{nm \frac{\rho - \rho'}{\rho} g}{s} = \frac{1}{3} \frac{mnV^2}{K},$$

or

$$V^2 = \frac{3Kg(\rho - \rho')}{s\rho}.$$

This equation enables  $V^2$  to be calculated from  $K$  and  $s$ . Perrin determined  $K$  by observing the positions of a number of the particles at known times and so getting the rate at which the mean square of their distances from a vertical plane increased with the time. This rate, as we have just seen, is equal to  $2K$ . He also found  $s$  by allowing a group of the particles to fall through the liquid for a known time. In this way he got an independent estimate of  $V^2$  and so of the ionic charge. This estimate agreed with his previous result got from the distribution of the particles with respect to height in the liquid.

Perrin's results provide an interesting confirmation of the atomic theory of matter and of electricity. His value of  $e$  is generally regarded as less reliable than that obtained by Millikan.

The value of the ionic charge was deduced by Planck from his theory of heat radiation. This is discussed in the chapter on the Quantum Theory.

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## CHAPTER XIV

# The Motion of Electrons in Gases

### 1. Townsend's Apparatus.

When ions are produced in gases, by X-rays or in other ways, at atmospheric pressure, it is found that they consist of clusters of several molecules carrying the electronic charge  $e$ . At lower pressures in many cases the positive ions are clusters of molecules, but the negative ions are merely free electrons. The motion of these electrons in gases has

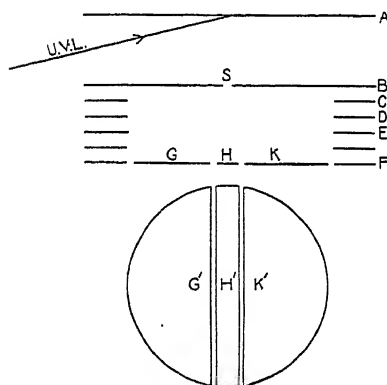


Fig. 1

been studied extensively by J. S. Townsend, of Oxford, and others, and this chapter is devoted mainly to a discussion of Townsend's experiments.

The apparatus used by Townsend is shown diagrammatically in fig. 1. A and B are two parallel circular metal plates about 4 cm. apart. At the centre of B there is a slit S which is 2 mm. wide and 1.5 cm. long. C, D, E, and F are metal rings cut out of thin sheets. These rings are equally spaced about 1 cm. apart. In the plane of the ring

F there is a circular disc GHK which nearly fills the hole in the ring. This disc is divided into three parts by two straight parallel cuts equally distant from the centre. The disc is shown in plan at G'H'K'. The different parts of the apparatus are supported inside a metal case not shown in the figure. The case can be evacuated and filled with gas at any desired pressure. By allowing ultra-violet light to fall on the under side of the plate A electrons can be set free at its surface. The plate B is kept at a higher potential than A so that the electrons emitted by A move through the gas towards B. Some of them pass through the slit S into the space between B and GHK. The disc GHK and the ring F are kept at a higher potential than the

plate B, so that the electrons which pass through the slit continue to move down and finally reach the disc. The rings C, D, E are used to make the electric field between S and H more nearly uniform. They are kept at potentials such that the potential differences between them and B are proportional to their distances from the plate B. The different potential differences are maintained by suitable batteries, and the disc GHK and ring F are kept at zero potential. The electric field above B is always made equal to that below, so that the electrons acquire a uniform velocity above B and continue to move with the same velocity below B. The length of the slit S is parallel to the sides of the strip H. The strip H is 4.5 mm. wide and 7 cm. long, with gaps on each side of it 0.5 mm. wide. The distance between the plate B and the disc GHK is 4 cm. As the electrons move down from the slit towards H they diffuse out sideways so that some of them fall on H and some on G and K. The three parts of the disc can be insulated and can be connected to a quadrant electrometer and induction balance by means of which the negative charges which they receive can be measured. Let  $n_1$ ,  $n_2$ , and  $n_3$  be the charges received by G, H, and K respectively. The sideways diffusion of the electrons then depends on the ratio

$$R = \frac{n_2}{n_1 + n_2 + n_3}. \quad \text{This ratio was determined for a number of gas}$$

pressures  $p$ , and values of the electric field strength  $Z$  along the path of the electrons.

## 2. Mathematical Theory of Townsend's Experiment.

Let us now consider the theory of this experiment. If  $K$  is the coefficient of diffusion of the electrons in the gas, then, using rectangular axes  $x$ ,  $y$ , and  $z$ , we have  $-K \frac{\partial n}{\partial x} = nv_x$ , where  $n$  is the number of electrons per unit volume, and  $v_x$  their average velocity component along  $x$ .

$$\text{Also} \quad -K \frac{\partial n}{\partial y} = nv_y,$$

$$-K \frac{\partial n}{\partial z} = nv_z.$$

The partial pressure  $P$  of the electrons is proportional to  $n$ , so that

$$-K \frac{\partial P}{\partial x} = Pv_x,$$

with similar equations in  $y$  and  $z$ . For a steady state, in which  $\frac{\partial n}{\partial t} = 0$  everywhere, we have  $\text{div}(n\mathbf{v}) = 0$ , or  $\text{div}(P\mathbf{v}) = 0$ , since  $n\mathbf{v}$  is the number of electrons flowing through unit area in unit time, and no new electrons are generated in the gas. The equation  $-\frac{\partial P}{\partial z} = \frac{Pv_z}{K}$  may be said to give the velocity of the electrons

due to a force  $-\frac{\partial P}{\partial z}$  on those in unit volume. If there is present an electric  $Z$  along the  $z$  axis, this gives an additional force  $neZ$  on the electrons, so

$$neZ - \frac{\partial P}{\partial z} = \frac{Pv_z}{K}.$$

In Townsend's apparatus take the origin at the centre of the slit and the  $z$ -vertically downwards along the path of the electrons. Also take  $y$  along slit, and  $x$  perpendicular to the slit in the plane of the plate B.

The equations of motion of the electrons are then:

$$\begin{aligned} -\frac{\partial P}{\partial x} &= \frac{Pv_x}{K}, \\ -\frac{\partial P}{\partial y} &= \frac{Pv_y}{K}, \\ neZ - \frac{\partial P}{\partial z} &= \frac{Pv_z}{K}. \end{aligned}$$

Differentiating these with respect to  $x, y, z$  respectively, and using  $\text{div}(P\mathbf{v})$ :

we get 
$$\frac{\partial^2 P}{\partial x^2} + \frac{\partial^2 P}{\partial y^2} + \frac{\partial^2 P}{\partial z^2} = Ze \frac{\partial n}{\partial z}.$$

The distribution of the electrons between the three parts of the disc is determined almost entirely by the diffusion along the  $z$  axis, so that for an approximate solution we may use  $\frac{\partial^2 P}{\partial z^2} = Ze \frac{\partial n}{\partial z}$ . This would be exactly correct if the were infinitely long. For a gas at temperature  $T$  we have  $P = nkT$ , where  $k$  is the gas constant for one molecule. It is found that the average kinetic energy of the electrons exceeds that of gas molecules, so that the temperature of electrons is greater than the temperature of the gas through which they moving. For the electrons, then, let  $P = \beta nkT$ , where  $T$  is the temperature of the gas and  $\beta T$  that of the electrons. This gives

$$\frac{\partial^2 n}{\partial x^2} = \frac{Ze}{\beta kT} \frac{\partial n}{\partial z}.$$

The factor  $e/kT$  is the same as  $Ne/NkT$ , where  $N$  is the number of molecules of one gramme-molecule of any gas. We have

$$\begin{aligned} Ne &= 9650 \text{ electromagnetic units,} \\ Nk &= 8.315 \times 10^7, \end{aligned}$$

so that, with  $T = 273 + 15$ , which was about the temperature in all Townsend's work, we get

$$\frac{\partial^2 n}{\partial x^2} = 40.3 \frac{Z}{\beta} \frac{\partial n}{\partial z},$$

where  $Z$  is now expressed in volts per centimetre. A solution of this equation

$$n = Az^{-1/2} e^{-\alpha x^2/z},$$

where  $A$  and  $\alpha$  are constants. This makes  $n = 0$  at  $z = 0$ , except where  $x = 0$ ; and so approximately satisfies the conditions in Townsend's apparatus; for

$z = 0$ ,  $n$  is zero, except on the slit where  $x$  is nearly zero. On substituting, we find  $\alpha = 10 \cdot 1Z/\beta$ . The ratio  $R$  is therefore given by

$$R = \frac{\int_{-b/2}^{+b/2} \epsilon^{-\alpha x^2/4z} dx}{\int_{-\infty}^{+\infty} \epsilon^{-\alpha x^2/4z} dx} = \frac{2}{\sqrt{\pi}} \int_0^{0.2\sqrt{Z/\beta}} \epsilon^{-y^2} dy,$$

since the breadth  $b$  of the strip  $H$ , plus half the total width of the gaps on each side of it, is 0.55 cm., and  $z = 4$  cm. at the disc. Townsend obtained a similar but more accurate solution of the problem, and so was able to find  $\beta$  from the experimentally found values of  $R$ . The above expression for  $R$  agrees nearly with Townsend's values.

It is found in this way that  $\beta$  is a function of  $Z/p$  in any given gas. That is, if the pressure  $p$  is changed and  $Z$ , the electric field strength, is also changed so as to keep  $Z/p$  the same, then  $\beta$  remains unaltered.

The following table gives some of the values of  $\beta$  (= temperature of electrons/temperature of gas—see above) found in nitrogen gas at 15° C.

$Z/p$		$\beta$
60	....	126.0
40	....	89.0
20	....	59.5
10	....	48.5
5	....	41.3
2	....	30.5
1	....	21.5
0.5	....	13.0
0.25	....	7.5

$Z$  is in volts per centimetre and  $p$  in millimetres of mercury.

Thus, for example, in an electric field of 20 volts per centimetre in nitrogen at 1 mm. pressure  $Z/p = 20$ , and so  $\beta = 59.5$ . That is, the kinetic energy of an electron is 59.5 times that of a gas molecule at 15° C. With  $Z = 100$  volts per centimetre and  $p = 5$  mm.,  $\beta$  is still 59.5.

### 3. Average Velocity and Kinetic Energy of the Electrons.

The average velocity of the electrons in the direction of the electric field  $Z$  was found with the same apparatus by deflecting the stream of electrons by means of a magnetic field. A uniform magnetic field of strength  $H$  was produced by means of suitable coils of wire through which a current could be passed. This field was perpendicular to the electric field  $Z$ , and parallel to the length of the slit. The stream of electrons was deflected sideways by the magnetic field, and the strength of the field was adjusted until one-half of the electrons was received by the electrode  $G$  and the other half by the electrodes  $H$  and  $K$  connected together. The centre of the stream was then deflected through 2.75 mm. at the plane  $GHK$ . By reversing the field the electrons could be deflected in the opposite direction so that half fell on  $K$  and half on  $G$  and  $H$  together. If the average velocity of the electrons along the

direction of the field  $Z$  is  $W$ , then the force on  $n$  electrons due to the magnetic field  $H$  is  $nHeW$  and the force due to the electric field is  $neZ$ . Consequently, if  $a$  is the deflection of the electrons while they move along the electric field a distance  $b$ , we have  $\frac{HW}{Z} = \frac{a}{b}$ . In Townsend's apparatus,  $a$  was 2.75 mm. when  $b$  was 40 mm., so that

$$W = \frac{Z}{H} \times 0.06875.$$

In this way  $W$  was found for different values of  $Z$  and  $p$ .

The average kinetic energy of the electrons is  $\beta$  times that of a gas molecule at the same temperature. The average energy of a gas molecule is  $\frac{3}{2}kT$ , where  $k$  is the gas constant for one molecule, and  $T$  is the absolute temperature. If  $m$  is the mass of one electron, and  $V^2$  the average of the squares of the velocities of the electrons, then

$$\frac{3}{2}\beta kT = \frac{1}{2}mV^2,$$

so that

$$V^2 = \frac{3\beta kT}{m}.$$

As the electrons move through the gas they acquire energy by the action of the electric field and lose it to the gas molecules by collisions. The velocity  $V$  with which the electrons move about is continually changed in direction by collisions, and it is much greater than  $W$ , the average velocity of drift along the electric field.  $V$  is usually called the velocity of agitation. It is easy to calculate  $W$  approximately. The rate at which  $n$  electrons receive momentum in the  $z$  direction owing to the electric field is  $neZ$ . The momentum of these electrons in the  $z$  direction when a steady state has been reached is  $nmW$ . When an electron collides with a gas molecule, we shall assume that after the collision it is as likely to be moving in one direction as another. Consequently the momentum of  $n$  electrons immediately after collisions is zero. On the average, then, an electron loses momentum  $mW$  per collision. The number of collisions made by  $n$  electrons in unit time is approximately  $nV/\lambda$ , where  $\lambda$  is the mean free path. Hence the momentum lost by  $n$  electrons in unit time is  $\frac{nV}{\lambda} \times mW$ , so that in a steady state we have

$$neZ = \frac{nmVW}{\lambda},$$

and therefore

$$\lambda = \frac{mVW}{Ze}.$$

Townsend's experiments give  $V$  and  $W$ , so that  $\lambda$  can be calculated.

The energy received by  $n$  electrons in unit time from the electric field is  $neZW$ , so that the average energy lost at a collision is  $\frac{neZW}{nV/\lambda}$ . Dividing this by  $\frac{1}{2}mV^2$ , we get the average fraction of the energy of the electron lost per collision. This is

$$\frac{2neZW\lambda}{nVmV^2} = \frac{2W^2}{V^2}, \text{ since } \frac{Ze\lambda}{mV} = W,$$

and so can be calculated.

The tables below give some of the results obtained in several gases, the symbols having the following meanings:  $Z$  = electric field;  $p$  = gas pressure;  $\beta$  = ratio of average energy of electron and gas molecule;  $W$  = average velocity of drift of electrons along electric field;  $V$  = their velocity of agitation (see above);  $f$  = ratio  $2W^2 : V^2$ .  $Z/p$  is in volts per centimetre per millimetre pressure,  $W$  and  $V$  are in centimetres per second, and  $\lambda$  is the mean free path in centimetres at 1 mm. pressure.

$Z/p$ .	$\beta$ .	$W$ .	$V$ .	$\lambda$ .	$f$ .
Nitrogen.					
60.0	126	$193 \times 10^5$	$129 \times 10^6$	0.0280	0.055
20.0	60	86 "	89 "	0.0266	0.0234
5.0	41	27 "	74 "	0.0277	0.0033
1.0	22	9 "	54 "	0.032	0.0007
0.25	8	5 "	32 "	0.045	0.0007
Hydrogen.					
50.0	148	$217 \times 10^5$	$140 \times 10^6$	0.042	0.059
20.0	78	70 "	102 "	0.025	0.0117
5.0	26	26 "	59 "	0.021	0.0046
1.0	9	12 "	35 "	0.029	0.0029
0.25	3	7 "	20 "	0.036	0.0026
Argon.					
15.0	324	$82 \times 10^5$	$207 \times 10^6$	0.079	0.000386
5.0	310	40 "	202 "	0.113	0.000097
0.95	280	6 "	193 "	0.085	0.000024
0.195	120	3 "	126 "	0.147	0.000016
Helium.					
5.0	172	$30 \times 10^5$	$151 \times 10^6$	0.064	0.00098
1.0	53	8.3 "	84 "	0.049	0.00024
0.1	6.2	3 "	29 "	0.06	0.00026
0.013	1.77	1.1 "	15 "	0.091	0.00013

The velocity of drift  $W$  is not proportional to  $Z/p$ ; it increases rather more rapidly than  $\sqrt{Z/p}$  in most cases. This is due mainly to the increase in the velocity

of agitation. In the inactive gases, argon and helium, the energy lost at collisions is very small, showing that the collisions are almost perfectly elastic; even in nitrogen the loss is only about one part in one thousand when  $Z/p$  is small.

The mean free path varies with  $Z/p$ , and seems to pass through a minimum value as  $Z/p$  increases. The mean free path in argon is surprisingly large when  $Z/p$  is small.

#### 4. Ionization by Collisions.

When a stream of electrons is passing through a gas in an electric field, and the kinetic energy of the electrons is great enough, some of the collisions between electrons and molecules cause an electron to escape from the molecule. A molecule is said to be ionized by such a collision, since a molecule which has lost an electron forms a positive ion. The ionization of gases by a stream of rapidly moving electrons was observed by Lenard, who passed Lenard rays, which are rapidly moving electrons, through different gases. The ionization of gases by collisions has been investigated by Townsend, and his results will be discussed here. Other important investigations on ionization by collisions are described in the chapter on ionization and radiation potentials. In Townsend's experiments electrons were set free at the surface of a metal plate by means of ultra-violet light. A uniform electric field was maintained between this plate and another parallel plate, which was charged positively, so that the electrons moved across from the first plate to the second. The charge received by the second plate was measured, and of course is proportional to the number of electrons which arrive at it. The space between the two plates was filled with a gas at pressure  $p$ , and the distance between the two plates was varied, keeping the electric field strength between them constant by making the potential difference proportional to the distance between the plates. It was found that the charge received by the second plate increased with the distance  $d$  between the plates, so that  $q = q_0 e^{\alpha d}$ , where  $q$  is the charge at distance  $d$ ,  $q_0$  the value of  $q$  when  $d$  is very small, and  $\alpha$  a constant.

Let us suppose that  $n_0$  electrons are set free by the ultra-violet light and that as these move across to the opposite plate they ionize the gas molecules, each setting free  $\alpha$  electrons from the gas molecules while moving 1 cm. in the direction of the electric field. Then if  $n$  electrons pass through a plane at a distance  $x$  from the plate from which  $n_0$  start, we have  $dn = \alpha n dx$ . This gives  $\log n = \alpha x + \text{constant}$ , so that since  $n = n_0$  at  $x = 0$  we get

$$n = n_0 e^{\alpha x}.$$

Thus, putting  $q = ne$  and  $q_0 = n_0 e$ , we get  $q = q_0 e^{\alpha d}$ , as was found experimentally. By measuring  $q$  and  $q_0$  Townsend determined the value of  $\alpha$  for different electric field strengths  $X$ , and different gas pressures  $p$ . It was found that  $\alpha/p$  is a function of  $X/p$  for a given

gas at a constant temperature, so that  $\alpha = pf(X/p)$ . This means, for example, that if  $X$  and  $p$  are both increased  $n$  times so that  $X/p$  remains the same, then  $\alpha$  is also increased  $n$  times. We have seen that the kinetic energy of the electrons is a function of  $X/p$ , and the same is true of the velocity  $W$  with which they move through the gas in the direction of the electric field. We should expect the number of molecules ionized to depend on the kinetic energy of the electrons, and to be proportional to the number of collisions between molecules and electrons. Suppose  $n$  electrons pass through a layer of the gas of thickness  $dx$ . The time each electron is in this layer is  $dx/W$ , and the  $n$  electrons make  $nV/\lambda$  collisions in unit time,  $V$  being the velocity of agitation and  $\lambda$  the mean free path, as before. The number of collisions in the layer is, therefore,

$$\frac{dx}{W} \frac{nV}{\lambda}.$$

If a fraction  $F$  of the collisions result in ionization, the number of electrons liberated per unit length is  $FnV/\lambda W$ , so that

$$\alpha = FV/\lambda W.$$

Now  $V$  and  $W$  are both functions of  $X/p$ ,  $\lambda$  is inversely as  $p$ , and  $F$  must depend on  $\frac{1}{2}mV^2$  or  $V$  only, and so is also a function of  $X/p$ , hence we must have  $\alpha = pf(X/p)$ , in agreement with the experimental results. If we suppose that all collisions for which the velocity of the electron is greater than a certain value result in ionization then we can calculate the fraction  $F$ . Let  $V$  now denote the velocity of agitation of an individual electron, and  $\bar{V}^2$  the average of the squares of the velocities of the electrons, and assume that the distribution of the velocities  $V$  is given by Maxwell's law; that is, let the number of electrons in unit volume for which  $V$  is between  $V$  and  $V + dV$  be  $AV^2e^{-qV^2}dV$ , so that the number of collisions made by these electrons in unit time is proportional to  $V^3e^{-qV^2}dV$ . The constant  $q$  will be equal to  $\frac{m}{2\beta kT}$ , where  $m$  is the mass of an electron and  $k$  is the gas constant for one molecule.

The number of collisions for which  $V$  is greater than  $V_0$ , divided by the total number, is therefore given by

$$F = \frac{\int_{V_0}^{\infty} V^3 e^{-qV^2} dV}{\int_0^{\infty} V^3 e^{-qV^2} dV},$$

Hence

$$F = (qV_0^2 + 1)e^{-qV_0^2},$$

so that

$$\alpha = \frac{\bar{V}}{\lambda W} (qV_0^2 + 1)e^{-qV_0^2}.$$

We have  $\lambda = \lambda_1/p$ , where  $\lambda_1$  is the mean free path at 1 mm. pressure, and also  $p = \frac{1}{3}mn\bar{V}^2 = \beta nkT$ , so that  $\beta kT = \frac{1}{3}m\bar{V}^2$ , and  $q = \frac{m}{2\beta kT}$ .

Hence

$$qV_0^2 = \frac{3}{2} \frac{V_0^2}{\bar{V}^2},$$

and

$$\alpha = \frac{p}{\lambda_1 W} \left( \frac{3}{2} \frac{V_0^2}{\bar{V}} + \bar{V} \right) e^{-3V_0^{3/2}/2\bar{V}^2}.$$

When  $\bar{V}/V_0$  is large, this becomes approximately  $\alpha = p\bar{V}/\lambda_1 W$ , so that if  $\bar{V}$  and  $W$  become equal when  $\bar{V}$  is large,  $\alpha = \frac{p}{\lambda_1} = \frac{1}{\lambda}$ . This theory indicates that  $\alpha$  should be very small when  $\bar{V}$  is much less than  $V_0$  and should increase rapidly with  $\bar{V}$  when  $V_0$  and  $\bar{V}$  are about equal. For large values of  $\bar{V}$  the rate of increase of  $\alpha$  with  $\bar{V}$  should become small. The assumption that all collisions for which  $V$  is greater than  $V_0$  result in ionization is not really correct, and it is found that the very fast electrons of cathode rays produce fewer ions than slower electrons, so that this theory is not satisfactory and is merely intended as a rough illustration. It is not possible to deduce from measurements of  $\alpha$  any accurate estimate of the velocity which an electron must have to be able to ionize a molecule. Accurate methods of measuring this velocity have been developed and are described in the chapter on ionization and radiation potentials.

The following table gives some of the values of  $\alpha/p$  found by Townsend in different gases for the values of  $X/p$  given in the first row of the table.

$X/p$ .	1000.	800.	600.	400.	200.	100.
Air .. ..	10.5	9.3	7.9	5.82	2.6	0.72
Water vapour	9.7	9.0	7.95	6.35	3.6	1.31
Hydrogen ..	—	—	—	3.7	2.62	1.36
Argon ..	—	—	9.2	7.5	4.4	2.0
Helium ..	—	—	—	—	2.37	2.0

## 5. Ionization by Positive Ions.

The equation  $q = q_0 e^{\alpha d}$ , which was found to give the charge  $q$  received by an electrode at a distance  $d$  from the plate at which electrons carrying a charge  $q_0$  start, gives  $q$  accurately when  $\alpha$  is small and  $d$  not too large. However, it was found, with large values of  $X/p$ , for which  $\alpha$  is large, that when  $d$  also was large the charge  $q$  obtained was greater than that given by this equation. This was explained by Townsend by means of the hypothesis that the positive ions also

ionize molecules by collisions when  $X/p$  is large. Another possible explanation is that the positive ions liberate electrons from the negatively charged electrode when they strike it. The results obtained can be explained equally well in either way, and Townsend's experiments do not enable us to decide which explanation is correct. We shall consider Townsend's explanation here, and the alternative one in section 7.

Let a positive ion ionize  $\beta$  molecules while it moves unit distance in the direction of the electric field  $X$  in gas at pressure  $p$ . Let  $n_0$  electrons start from an electrode at  $x = 0$ , and let  $n$  be the number which pass through a plane at a distance  $x$  from this electrode. Also let  $m$  be the number of positive ions which pass through this plane in the opposite direction. Then between the planes at  $x$  and  $x + dx$  the number of molecules ionized is

$$n\alpha dx + m\beta dx,$$

so that

$$dn = n\alpha dx + m\beta dx,$$

and

$$-dm = n\alpha dx + m\beta dx,$$

since each molecule ionized gives an electron and a positive ion. This gives  $dn + dm = 0$ , so that  $n + m = \text{constant}$ . At  $x = 0$  we have  $n = n_0$ , and if  $d$  is the distance between the electrodes then at  $x = d$  we have  $m = 0$ , so that the constant is equal to  $n_d$ . Hence

$$n + m = n_d,$$

and

$$dn = n\alpha dx + (n_d - n)\beta dx = n(\alpha - \beta)dx + n_d\beta dx.$$

Integrating this, we get

$$\frac{1}{\alpha - \beta} \log \{n(\alpha - \beta) + n_d\beta\} = x + \text{constant}.$$

At  $x = 0$ ,  $n = n_0$ , so that

$$\frac{1}{\alpha - \beta} \log \{n_0(\alpha - \beta) + n_d\beta\} = \text{constant}.$$

Hence

$$\frac{1}{\alpha - \beta} \log \frac{n(\alpha - \beta) + n_d\beta}{n_0(\alpha - \beta) + n_d\beta} = x.$$

At  $x = d$ ,  $n = n_d$ , so that

$$\frac{1}{\alpha - \beta} \log \frac{n_d\alpha}{n_0(\alpha - \beta) + n_d\beta} = d,$$

or

$$n_d = \frac{n_0(\alpha - \beta)}{\alpha e^{-d(\alpha - \beta)} - \beta}.$$

When  $\beta = 0$ , this reduces to

$$n_d = n_0 \epsilon^{\alpha d},$$

as we should expect. By measuring  $q = n_d e$  and  $q_0 = n_0 e$  for different values of  $d$ , the values of  $\alpha$  and  $\beta$  can be determined. When  $d$  is small,  $n_d = n_0 \epsilon^{\alpha d}$  nearly, which gives an approximate value of  $\alpha$ . The best values of  $\alpha$  and  $\beta$  can then be found by trial.

The following table contains some results given by Townsend for air at 4 mm. pressure with an electric field of 700 volts per centimetre. The value of  $\alpha$  used was 8.16 and that of  $\beta$  was 0.0067.

$d$ .	$q/q_0$ .	$\epsilon^{\alpha d}$ .	$\frac{\alpha - \beta}{\alpha \epsilon^{-d(\alpha - \beta)} - \beta}$ .
Cm			
0.2	5.12	4.9	5.11
0.3	11.4	11.6	11.6
0.4	26.7	26.1	26.5
0.5	61.0	59.0	62.0
0.6	148.0	133.0	149.0
0.7	401.0	301.0	399.0
0.8	1500.0	680.0	1544.0

We see that  $\epsilon^{\alpha d}$  represents the values of  $q/q_0$  quite well up to  $d = 0.5$  cm., but for larger values of  $d$ ,  $\epsilon^{\alpha d}$  is too small. The expression  $\frac{\alpha - \beta}{\alpha \epsilon^{-d(\alpha - \beta)} - \beta}$ , however, agrees well with all the values of  $q$ . Similar results were obtained by Townsend with air and other gases at different pressures, and with different electric fields. It was found that  $\beta = p \phi\left(\frac{X}{p}\right)$ , where  $\phi\left(\frac{X}{p}\right)$  denotes a function of  $X/p$ .  $\beta$  is always much smaller than  $\alpha$ . The following table gives some values of  $\beta/p$ .

$X/p$ .	Hydrogen.	Argon.	Air.	Carbon Dioxide.
200	0.08	0.02	—	—
600	—	0.22	0.10	—
800	—	—	0.18	0.01
1400	—	—	—	0.10

## 6. Sparking Potentials.

The equation

$$\frac{q}{q_0} = \frac{\alpha - \beta}{\alpha \epsilon^{-d(\alpha - \beta)} - \beta}$$

shows that  $q/q_0$  becomes infinite when  $\alpha = \beta \epsilon^{d(\alpha - \beta)}$ . This means that,

a discharge once started would continue indefinitely. The equation  $\alpha = \beta e^{d(\alpha-\beta)}$  therefore gives the sparking potential for the distance  $d$ , pressure  $p$ , and field strength  $X$ . The sparking potential  $S$  is given by  $S = Xd$ . When  $q/q_0$  is infinite a single electron set free at the negative electrode is sufficient to start a continuous electric discharge. Now the gas always contains a few free electrons owing to the presence of traces of radioactive bodies, so that if the potential difference between the two parallel plates is gradually increased until the field reaches the value for which  $\alpha = \beta e^{d(\alpha-\beta)}$ , a continuous discharge will then start in the gas between the plates.

Now  $\frac{X'}{p} = \frac{S}{pd}$ , where  $X'$  is the field strength for the sparking potential  $S$ . Hence we have  $\alpha = pf(S/pd)$ , and  $\beta = p\phi(S/pd)$ , where  $f(S/pd)$  and  $\phi(S/pd)$  denote functions of  $S/pd$ . Hence the equation  $\alpha = \beta e^{d(\alpha-\beta)}$  may be written

$$f(S/pd) = \phi(S/pd) e^{pd[f(S/pd) - \phi(S/pd)]}.$$

This is a relation between  $S$  and  $pd$ , so that we must have  $S = S(pd)$ , where  $S(pd)$  denotes a function of  $pd$  only. It is found experimentally that the sparking potential between parallel plates in any gas depends on the product of the gas pressure and the distance between the plates. Townsend found that the observed sparking potentials agreed well with those given by the equation

$$\alpha = \beta e^{d(\alpha-\beta)}.$$

For example, in air at 4 mm. pressure, with an electric field of 700 volts per centimetre, it was found that  $\alpha = 8.16$  and  $\beta = 0.0067$ . These values of  $\alpha$  and  $\beta$  give  $d = 0.871$  cm. when substituted in the condition for sparking. Hence  $S = 0.871 \times 700 = 609$  volts. It was found that with  $d = 0.871$  cm. a continuous discharge began when the potential was increased to 615 volts, which agrees well with 609 volts.

The sparking potential  $S = S(pd)$  has a minimum for a certain value of  $pd$ . If  $p$  is kept constant and  $d$  gradually increased from a very small value,  $S$  at first falls as  $d$  increases, then passes through a minimum value and rises again. When  $pd$  is large,  $S$  is nearly proportional to  $pd$ . If  $d$  is kept constant and the gas pressure varied, when  $p$  is very small the sparking potential is very large. As  $p$  increases the sparking potential falls to the minimum value, and then rises and becomes nearly proportional to  $p$ .

The following table gives some sparking potentials  $S$  in several gases between parallel plates 3 mm. apart. The pressures  $p$  are in millimetres of mercury, and  $S$  is in volts.

Air.		Hydrogen.		Sulphur Dioxide.	
<i>p.</i>	<i>S.</i>	<i>p.</i>	<i>S.</i>	<i>p.</i>	<i>S.</i>
51.0	1480	13.6	415	13.5	1145
21.4	790	8.54	356	4.5	651
5.99	452	5.4	301	1.61	471
2.51	371	4.02	278	1.04	457
1.89	356	3.44	282	0.8	465
1.22	375	2.52	310	0.43	621
0.928	441	2.15	356	0.23	1590
0.536	863	1.35	780	—	—
0.357	1786	0.861	1789	—	—

These results are due to Carr. The two parallel electrodes were separated by a ring of ebonite 3 mm. thick, which prevented the discharge passing between the backs or the edges of the electrodes. This precaution is necessary when the pressure is below that at which *S* is a minimum, because then the discharge tends to pass across any available path which is longer than the shortest distance between the electrodes. The fact that in any gas the sparking potential is a function of *pd* was first discovered by Paschen.

### 7. Alternative Theory of Action of Positive Ions.

Let us now suppose that the positive ions do not ionize the gas molecules, but that they set free electrons when they strike the negative electrode. Let  $n_0$  electrons set free by ultra-violet light start from the negative electrode and move across to the positive electrode. The number of electrons which arrive at the positive electrode is then  $n_0\epsilon^{ad}$ , and the number of positive ions produced is  $n_0(\epsilon^{ad} - 1)$ . These positive ions strike the negative electrode, and we suppose that each on the average sets free  $\gamma$  electrons at it. Hence  $\gamma n_0(\epsilon^{ad} - 1)$  are set free and move across like the original  $n_0$  electrons, and so on. The total number of electrons that finally get to the positive electrode is therefore

$$n = n_0\epsilon^{ad} \{1 + \gamma(\epsilon^{ad} - 1) + \gamma^2(\epsilon^{ad} - 1)^2 + \dots\}$$

$$= \frac{n_0\epsilon^{ad}}{1 - \gamma(\epsilon^{ad} - 1)}$$

If we put  $\gamma = \frac{\beta'}{\alpha - \beta'}$ , we get

$$n = \frac{n_0(\alpha - \beta')\epsilon^{ad}}{\alpha - \beta'\epsilon^{ad}}.$$

Comparing this with the equation previously obtained on the first theory, viz.

$$n = \frac{n_0(\alpha - \beta)\epsilon^{d(\alpha - \beta)}}{\alpha - \beta\epsilon^{d(\alpha - \beta)}},$$

we see that since  $\beta$  is always very small compared with  $\alpha$  the two equations are practically identical. It follows that Townsend's experiments may be explained equally well by supposing that the positive ions only act by liberating electrons at the negative electrode. It has been shown experimentally that positive ions do set free electrons when they strike a metallic surface in a vacuum, but further experiments are necessary to decide on the relative importance of the two effects under different conditions. Some physicists consider that positive ions do not ionize molecules by collisions under any circumstances.

## REFERENCE

*Electricity in Gases.* J. S. Townsend.

## CHAPTER XV

# The Electrical Conductivity of Flames

### 1. Conductivity of a Bunsen Flame.

The fact that flames and the gases coming from them conduct electricity was known to Faraday and has been the subject of many investigations since his time. The ionic theory of conductivity was

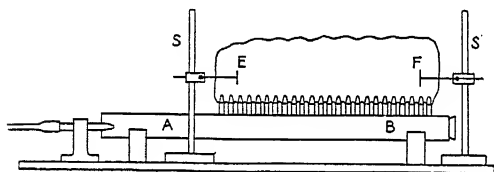


Fig. 1

first applied to gases by Giese as an explanation of his experiments on the conductivity of flame gases.

The electrical conductivity of a Bunsen flame can be conveniently studied with the

apparatus shown in fig. 1. A brass tube AB is supported horizontally on a wooden stand. The end near B is closed and coal gas is passed into the other end of the tube from a jet as shown. Small quartz tubes are cemented into the brass tube in a row about 25 cm. long, and the mixture of gas and air formed in the tube is burned at the ends of these quartz tubes. In this way a Bunsen flame about 10 cm. high and 26 cm. long is obtained. The quartz tubes serve to insulate the flame, so that its conductivity can be measured between two platinum electrodes E and F supported by movable stands S and S'. The electrodes may be made of pieces of sheet platinum about  $1.5 \times 1.5$  cm. welded to stout platinum wires. If the electrodes are connected to a battery through a galvanometer or micro-ammeter the current through the flame can be measured and the way it varies with the potential difference and distance between the electrodes observed.

It is found that the relation between the current  $C$ , the potential difference  $V$ , and the distance  $d$  between the electrodes is approximately

$$V = ACd + BC^2,$$

where  $A$  and  $B$  are constants, for any particular flame and electrodes. When  $d$  is small, say 1 or 2 mm., the term  $ACd$  is negligible, and the

current is nearly proportional to the square root of the potential difference.

The following table gives some values of the current in amperes obtained with a Bunsen flame similar to that just described.

Potential Difference.	Distance between Electrodes.		
	1 Cm.	9 Cm.	18 Cm.
600 volts	$310 \times 10^{-8}$	$295 \times 10^{-8}$	$270 \times 10^{-8}$
200 "	175 "	165 "	143 "
40 "	67 "	57 "	48 "
10 "	22 "	16 "	13 "
2 "	5 "	4 "	3 "

If one of the electrodes is moved near to the surface of the flame so that it becomes cooler the current is decreased. This effect is much more marked with the negative electrode than with the positive electrode.

## 2. Potential Differences in the Flame.

The difference of potential between any point in the flame and one of the electrodes can easily be measured by putting in a fine insulated platinum wire and connecting it and the electrode to an electrostatic voltmeter or quadrant electrometer. The instrument used must be insulated so that the potential in the flame may not be disturbed by it. The platinum probe wire can be covered with a small fused quartz tube about  $\frac{1}{2}$  mm. in diameter except for one or two millimetres at the end of the wire. In this way it is found that there is a nearly uniform potential gradient in the flame except near the electrodes. Close to the negative electrode there is usually a large and sudden drop of potential and a similar but much smaller drop close to the positive electrode. This is so when both electrodes are red hot and not near the surface of the flame. If either electrode is cooled by moving it near to the surface of the flame the drop of potential near it increases, and becomes nearly equal to the potential difference between the two electrodes if the electrode is allowed to get much cooler than the other one. The effect of cooling the negative electrode slightly is much greater than that of cooling the positive electrode.

The uniform potential gradient is approximately proportional to the current. The potential difference between the electrodes may therefore be regarded as made up of three parts  $V_1$ ,  $V_2$ , and  $V_3$ .  $V_1$  is the potential drop at the positive electrode,  $V_2$  that at the negative electrode, and  $V_3$  that due to the uniform gradient.  $V_3 = ACd$ , where  $A$  is a constant, so that

$$V = ACd + V_1 + V_2.$$

Comparing this with the equation  $V = ACd + BC^2$ , we see that  $V_1 + V_2 = BC^2$ , and this result can be easily verified by measuring  $V_1$  and  $V_2$  with different currents passing through the flame. The layers near the electrodes in which the potential drops occur are thicker at the negative electrode than at the positive electrode. The layer at the negative electrode may be several millimetres thick while that at the positive electrode is less than 1 mm. when both electrodes are red hot.

The temperature of the hottest part of a Bunsen flame is about  $2000^\circ \text{K}$ . By altering the proportion of gas to air used, the temperature can be varied, and by mixing an inert gas like carbon dioxide or nitrogen with the air supplied to the flame the temperature can be considerably reduced without putting the flame out. It is found that the ratio of the current to the uniform potential gradient in the flame, which may be taken as a measure of the conductivity, increases rapidly with the temperature.

### 3. Ions and Electrons: Theory of Conductivity of Flames.

An ordinary Bunsen flame is a mixture of nitrogen, water vapour, carbon monoxide, and dioxide, with some hydrogen, methane, and other hydrocarbons. It is supposed that some of the molecules present in the flame become ionized by collisions with each other or with electrons or by the action of light radiation; that is, electrons are set free from some molecules so that we get positive ions and electrons in the flame. Some of the electrons may get attached to molecules, so forming negative ions. It is probable that most of the electrons set free do not get attached to molecules, at any rate in the hotter parts of the flame.

The conductivity of the flame is due to the presence of these ions and electrons. Let  $e$  denote the charge on a positive ion so that  $-e$  is the charge on one electron, and let there be  $n_1$  positive ions and  $n_2$  electrons present in unit volume. Then, when there is an electric field in the flame, the positive ions drift along in the direction of the field and the electrons in the opposite direction. If  $i$  denotes the current density, or current per unit area of cross-section, we have

$$i = e(n_1 v_1 + n_2 v_2),$$

where  $v_1$  is the velocity of drift of the positive ions, and  $v_2$  that of the electrons. The velocities  $v_1$  and  $v_2$  are probably nearly proportional to the field strength  $X$ , at any rate when  $X$  is small. Hence  $v_1 = k_1 X$  and  $v_2 = k_2 X$ , where  $k_1$  and  $k_2$  are the velocities due to unit field and are usually called the mobilities of the ions and electrons respectively. We have then

$$i = eX(n_1 k_1 + n_2 k_2).$$

The volume density of charge in the flame is equal to  $e(n_1 - n_2)$ , so that we have

$$\frac{dX}{dx} = 4\pi e(n_1 - n_2),$$

where  $x$  is the distance of the point considered from the positive electrode. In

the uniform gradient between the electrodes  $\frac{dX}{dx} = 0$ , so that  $n_1 = n_2$  and therefore  $i = neX_0(k_1 + k_2)$ , where  $n = n_1 = n_2$  and  $X_0$  is the value of  $X$  in the uniform gradient. Now  $V_3$ , the fall of potential due to the uniform gradient, is equal to  $X_0d$ , so that, since  $V_3 = ACd$ , we get  $X_0 = AC$ . If  $S$  is the area of cross-section of the flame, then, if we assume that the current is uniformly distributed over the cross-section, we have  $i = C/S$ , and

$$A = \frac{1}{Sne(k_1 + k_2)}.$$

Also  $k_2$  is much larger than  $k_1$ , so that approximately  $A = 1/Snek_2$ .

The fact that the current is found to be nearly proportional to the uniform potential gradient shows therefore that the velocity of the electrons is nearly proportional to the field strength. This is true for fields up to 20 or 30 volts per centimetre. Near the electrodes, where the potential drops occur,  $dX/dx$  is not zero, so that  $n_1$  is not equal to  $n_2$ .

The positive ions and the electrons attract each other so that the electrons tend to recombine with positive ions, so reforming neutral molecules. It is easy to see that this recombination must be proportional to both  $n_1$  and  $n_2$  or to  $\alpha n_1 n_2$ , where  $\alpha$  is a constant usually called the coefficient of recombination.  $\alpha n_1 n_2$  is equal to the number of neutral molecules formed per unit volume in unit time by the combination of electrons and positive ions.

Consider a layer of thickness  $dx$  between the planes at  $x$  and  $x + dx$ . The number of positive ions flowing into this layer per unit area per unit time through the plane at  $x$  is  $n_1 v_1$ , and the number flowing out through the plane at  $x + dx$  is  $n_1 v_1 + \frac{d}{dx}(n_1 v_1)dx$ . The number which disappear by recombination in the layer is  $\alpha n_1 n_2 dx$ , so that if  $q$  is the number of molecules ionized per unit volume per unit time, which we shall suppose constant from one electrode to the other, then

$$\frac{d}{dx}(n_1 v_1) = q - \alpha n_1 n_2.$$

In the same way, for the electrons,

$$-\frac{d}{dx}(n_2 v_2) = q - \alpha n_1 n_2.$$

In the uniform gradient

$$\frac{d}{dx}(n_1 v_1) = \frac{d}{dx}(n_2 v_2) = 0,$$

so that  $q = \alpha n^2$ , or the ionization is equal to the recombination.

If no ions are emitted by the positive electrode, then  $n_1 = 0$  at its surface, that is at  $x = 0$ . Thus in the layer at the surface of the positive electrode in which  $X$  is greater than  $X_0$ , the flow of positive ions increases from zero at  $x = 0$  to  $nk_1 X_0$ , its value in the uniform gradient  $X_0$ . Let the thickness of this layer be  $\lambda_1$ ; then we have

$$\int_0^{\lambda_1} \frac{d}{dx}(n_1 v_1) dx = nk_1 X_0 = \int_0^{\lambda_1} (q - \alpha n_1 n_2) dx.$$

This equation merely expresses the fact that in the layer  $\lambda_1$  at the positive electrode there is an excess of ionization over recombination sufficient to supply the positive ions which flow across the uniform gradient. In the same way for the

layer near the negative electrode, if this electrode emits no electrons,  $n_2 = 0$  at its surface, and

$$-\int_{d-\lambda_2}^d \frac{d}{dx}(n_2 v_2) dx = nk_2 X_0 = \int_{d-\lambda_2}^d (q - \alpha n_1 n_2) dx,$$

where  $d$  is the distance between the electrodes, and  $\lambda_2$  the thickness of the layer at the negative electrode, in which  $q > \alpha n_1 n_2$ .

Now  $i/e = n_1 v_1 + n_2 v_2$ , so that  $n_1 v_1 + n_2 v_2$  is constant from one electrode to the other. Thus  $n_1 v_1 e$  increases from 0 at  $x = 0$  to  $enk_1 X_0$  at  $x = \lambda_1$ , remains constant from  $x = \lambda_1$  to  $d - \lambda_2$ , and increases from  $enk_1 X_0$  at  $x = d - \lambda_2$  to  $i$  at  $x = d$ ; while  $n_2 v_2 e$  is equal to  $i$  at  $x = 0$ , decreases to  $enk_2 X_0$  at  $x = \lambda_1$ , remains constant from  $x = \lambda_1$  to  $x = d - \lambda_2$ , and decreases from  $enk_2 X_0$  at  $x = d - \lambda_2$  to zero at  $x = d$ . Dividing  $nk_2 X_0$  by  $nk_1 X_0$  we get

$$\frac{k_2}{k_1} = \frac{\int_{d-\lambda_2}^d (q - \alpha n_1 n_2) dx}{\int_0^{\lambda_1} (q - \alpha n_1 n_2) dx}.$$

Now  $q - \alpha n_1 n_2$  is equal to  $q$  at  $x = 0$  and to zero at  $x = \lambda_1$ , and it is equal to zero at  $x = d - \lambda_2$  and to  $q$  at  $x = d$ , so that the average value of  $q - \alpha n_1 n_2$  over  $\lambda_1$  cannot differ much from that over  $\lambda_2$ . Hence the ratio of the two integrals must be nearly equal to  $\lambda_2/\lambda_1$ , so that we get  $k_2/k_1 = \lambda_2/\lambda_1$  approximately.

When both electrodes are red hot it is found that  $\lambda_2$  is much greater than  $\lambda_1$ , so that  $k_2$  must be greater than  $k_1$ .  $\lambda_1$  is so small that it cannot be determined accurately, whereas  $\lambda_2$  is several millimetres. We should expect the electrons to have a much greater velocity than positive ions, since the mass of an electron is several thousand times smaller than that of an ion.

The drops of potential in the layers  $\lambda_1$  and  $\lambda_2$  can be calculated approximately as follows:

The equation 
$$\frac{dX}{dx} = 4\pi e(n_1 - n_2),$$

multiplied by

$$X = \frac{v_1}{k_1} = \frac{v_2}{k_2},$$

gives

$$\frac{dX^2}{dx} = 8\pi e \left( \frac{n_1 v_1}{k_1} - \frac{n_2 v_2}{k_2} \right).$$

Differentiating this with respect to  $x$ , and substituting for  $\frac{d}{dx}(n_1 v_1)$  and  $-\frac{d}{dx}(n_2 v_2)$  the value  $q - \alpha n_1 n_2$ , we get

$$\frac{d^2 X^2}{dx^2} = 8\pi e \left( \frac{1}{k_1} + \frac{1}{k_2} \right) (q - \alpha n_1 n_2).$$

Now at  $x = 0$  we have  $n_1 = 0$ , and at  $x = \lambda_1$ ,  $q = \alpha n_1 n_2 = \alpha n^2$ , so that  $q - \alpha n_1 n_2$  changes from  $q$  at  $x = 0$  to 0 at  $x = \lambda_1$ . For an approximate calculation we may therefore take  $q - \alpha n_1 n_2 = q \left( 1 - \frac{x}{\lambda_1} \right)$  for values of  $x$  between 0 and  $\lambda_1$ . Assuming this, we get

$$\frac{d^2 X^2}{dx^2} = 8\pi e \left( \frac{1}{k_1} + \frac{1}{k_2} \right) q \left( 1 - \frac{x}{\lambda_1} \right),$$

and by integration 
$$\frac{dX^2}{dx} = 8\pi e \left( \frac{1}{k_1} + \frac{1}{k_2} \right) q \left( x - \frac{x^2}{2\lambda_1} \right) + \text{constant}.$$

The equation  $\frac{dX}{dx} = 4\pi e(n_1 - n_2)$  at  $x = 0$  becomes  $\frac{dX}{dx} = -4\pi en_2$ , and also at  $x = 0$  we have  $i = n_2 ek_2 X$ , so that

$$\left. X \frac{dX}{dx} \right|_{x=0} = -\frac{4\pi i}{k_2},$$

or

$$\left. \frac{dX^2}{dx} \right|_{x=0} = -\frac{8\pi i}{k_2}.$$

Using this to determine the constant we get

$$\frac{dX^2}{dx} = 8\pi e \left( \frac{1}{k_1} + \frac{1}{k_2} \right) q \left( x - \frac{x^2}{2\lambda_1} \right) - \frac{8\pi i}{k_2}.$$

At  $x = \lambda_1$ ,  $X$  becomes constant so that  $\frac{dX^2}{dx} = 0$ , therefore

$$\lambda_1 = \frac{2i}{qe} \frac{k_1}{k_1 + k_2},$$

so that

$$\frac{dX^2}{dx} = -\frac{8\pi i}{k_2} \left\{ 1 - \frac{x}{\lambda_1} \right\}^2.$$

Integrating this and putting  $X^2 = X_0^2$  at  $x = \lambda_1$

we get 
$$X^2 = \frac{8\pi i \lambda_1}{3k_2} \left( 1 - \frac{x}{\lambda_1} \right)^3 + X_0^2.$$

Since  $X$  is large compared with  $X_0$  except when  $X$  is nearly equal to  $\lambda_1$ , we have approximately from this:

$$X = \sqrt{\frac{8\pi i \lambda_1}{3k_2} \left( 1 - \frac{x}{\lambda_1} \right)^3}.$$

Hence the drop of potential in the layer  $\lambda_1$  is given by

$$V_1 = \int_0^{\lambda_1} \left( \frac{8\pi i \lambda_1}{3k_2} \right)^{\frac{1}{2}} \left( 1 - \frac{x}{\lambda_1} \right)^{3/2} dx.$$

This gives

$$V_1 = \left( \frac{8\pi i}{3k_2} \right)^{\frac{1}{2}} \frac{2}{5} \lambda_1^{3/2}.$$

Now

$$\lambda_1 = \frac{2ik_1}{qe(k_1 + k_2)},$$

so that finally

$$V_1 = \frac{1}{5} \left( \frac{\pi}{3k_2} \right)^{1/2} \left( \frac{k_1}{qe(k_1 + k_2)} \right)^{3/2} i^2.$$

In the same way we obtain for the potential drop  $V_2$  at the negative electrode

$$V_2 = \frac{1}{5} \left( \frac{\pi}{3k_1} \right)^{1/2} \left( \frac{k_2}{qe(k_1 + k_2)} \right)^{3/2} i^2.$$

Hence approximately

$$\frac{V_2}{V_1} = \left(\frac{k_2}{k_1}\right)^{1/2} \left(\frac{k_2}{k_1}\right)^{3/2} = \left(\frac{k_2}{k_1}\right)^2,$$

The difference of potential between the electrodes  $V$  is therefore given by

$$V = \frac{id}{ne(k_1 + k_2)} + \frac{1}{5} \left(\frac{\pi}{3k_1 k_2}\right)^{1/2} \left(\frac{1}{qe(k_1 + k_2)}\right)^{3/2} (k_1^2 + k_2^2) i^2.$$

This equation agrees with that found experimentally, viz:

$$V = ACd + BC^2.$$

The ratio  $V_2/V_1$  cannot be obtained accurately from the observed distribution of the potential between the electrodes when both electrodes are red hot, because  $V_1$  is too small to measure accurately, but it is clear that the ratio is quite large. The electrodes cool the flame so that the ionization close to the electrodes must be less than elsewhere in the flame, whereas we have supposed  $q$  constant over the distance between the electrodes.

This approximate theory of the relation between the current and the potential in the flame agrees in a general way with the facts, so that we may say that the results obtained are in accordance with the ionic theory. The two layers  $\lambda_1$  and  $\lambda_2$  have a total thickness

$$\lambda_1 + \lambda_2 = \frac{2i}{qe},$$

and we have supposed that there is a uniform potential gradient between the layers. As the potential difference increases  $i$  increases, so that  $\lambda_1 + \lambda_2$  also increases and so would eventually become equal to  $d$ , the distance between the electrodes. The current would then be equal to  $\frac{1}{2}qed$ , which is one-half the saturation current  $qed$  which would be obtained if there were no recombination. The theory therefore is only applicable when the current density is less than one-half the saturation current density.

When the potential difference between red-hot electrodes in a uniform flame is gradually increased, then at first the current is given approximately by the equation

$$V = ACd + BC^2,$$

but, when  $V$  becomes equal to a certain large value which depends on the distance between the electrodes, the current begins to increase much more rapidly with  $V$  than the above equation indicates. If  $V$  is increased much farther an arc discharge starts and the electrodes are melted. This rapid increase of the current is due to ionization by collisions. Ionization by collisions in gases at ordinary temperatures is discussed in the chapter on the motion of electrons in gases, and it obeys the same laws in flame and so need not be further discussed here. The currents obtained before ionization by collisions sets in are probably very small compared with the saturation current.

#### 4. Electron Mobilities in Flames.

We have supposed that the velocity  $v_2$  of the electrons along the direction of the electric field  $X$  is given by  $v_2 = k_2 X$ . According to the theory of the motion of electrons in gases we have approximately  $k_2 = e\lambda/m\bar{V}$ , where  $\lambda$  is the mean free path of the electrons,  $e$  the charge and  $m$  the mass of an electron, and  $\bar{V}$  the average velocity of agitation

of the electrons. Townsend's experiments show that in gases at ordinary temperatures  $\bar{V}$  is a function of  $X/p$ , where  $p$  is the gas pressure, and that the average kinetic energy of agitation of the electrons  $\frac{1}{2}m\bar{V}^2$  is much greater than that of a gas molecule at the same temperature, provided  $X/p$  is not extremely small. For example, in nitrogen with  $X/p$  equal to 0.25,  $X$  being in volts per centimetre and  $p$  in millimetres of mercury, he found  $\bar{V}$  to be 7.5 times greater than the value corresponding to the average energy of a gas molecule.

In a flame at about 2000° K. the gas density is about  $\frac{1}{7}$  of that at the ordinary temperature, and so is the same as for a gas at the ordinary temperature at about 100 mm. pressure. Thus in a flame with an electric field of 25 volts per centimetre we should expect the velocity of the electrons in the direction of the field to be about the same as Townsend found in nitrogen with  $X/p = 0.25$ , which was  $5 \times 10^5$  cm. per second. This makes  $k_2 = 5 \times 10^5 / 25 = 2 \times 10^4$  cm./sec. for 1 volt per centimetre. This estimate is probably too high because the kinetic energy of the electrons in the flame with  $X = 0$  is seven times that at the ordinary temperature. If we suppose that the field increases the kinetic energy in the same ratio at any temperature, then with  $X = 25$  volts/cm. we get

$$k_2 = 2 \times 10^4 / \sqrt{7} = 7500 \frac{\text{cm.}}{\text{sec.}} \text{ per } \frac{\text{volt}}{\text{cm.}}$$

Various attempts to estimate  $k_2$  in flames, by finding the field required to make the electrons move down the flame against the upward stream of gases, and by other similar methods, have been made and results varying from 1000  $\frac{\text{cm.}}{\text{sec.}}$  to 30,000  $\frac{\text{cm.}}{\text{sec.}}$  for 1 volt per centimetre obtained. Such methods are very difficult in flames, because the gases are strongly ionized throughout the flame so that it is difficult to prove that electrons are moving down the flame. Estimates of  $k_2$  by indirect methods are therefore more reliable.

Probably the best method of getting the mobility of the electrons in a Bunsen flame is by measuring the Hall Effect. A flame like that described at the beginning of this chapter is placed between the poles of a large electromagnet, which when excited produces a horizontal field perpendicular to the plane of the flame. Two fine platinum wires are mounted on the end of a shaft which turns in a hole bored in one of the poles of the magnet parallel to the direction of the magnetic field. The two wires are parallel to the magnetic field and about 1 cm. apart, the axis of rotation of the shaft being half-way between them. The ends of the wires project into the flame, and they are insulated and connected to an insulated quadrant electrometer. When a current is passed horizontally between electrodes in the flame so that the

two wires are in the uniform potential gradient  $X_0$  between the electrodes, the quadrant electrometer will indicate a potential difference equal to  $X_0 l \sin \theta$ , where  $l$  is the distance between the two wires, and  $\theta$  is the angle between the plane containing the two wires and a plane perpendicular to the electric field  $X_0$ . By turning the shaft the angle  $\theta$  can be varied so that it is easy to make the potential difference zero, in which case  $\theta = 0$ . If now the magnet is excited the quadrant electrometer is deflected, but by rotating the shaft the deflection can be brought back to zero. The magnetic field rotates the equipotential planes in the flame through an angle  $\phi$  which can be measured in this way.

If  $v_2$  is the velocity of the electrons due to the electric field  $X_0$ , then the magnetic field will give rise to a vertical force on them equal to  $Hev_2$ . Since the flame is insulated, this force cannot produce a vertical current, so that a vertical electric field is produced of strength  $Z$  such that

$$Ze = Hev_2.$$

The equipotential planes are therefore turned through the angle  $\phi$  given by

$$\tan \phi = \frac{Z}{X_0},$$

so that

$$v_2 = \frac{Z}{H} = \frac{X_0}{H} \tan \phi.$$

If we put  $v_2 = k_2 X_0$ , this becomes

$$k_2 = \frac{\tan \phi}{H}.$$

In this way it has been found that  $k_2$  is about 4000  $\frac{\text{cm.}}{\text{sec.}}$  for 1 volt per centimetre. This high mobility of the negative ions in flames shows that they must be electrons.

The conductivity of the flame as measured by the ratio of the current to the uniform field  $X_0$  is found to be somewhat diminished by a transverse magnetic field. When the current is horizontal, and the magnetic field is also horizontal and perpendicular to the current, there is a vertical mechanical force on the flame, as with any conductor carrying a current in a magnetic field. This force retards the upward motion of the flame when the magnetic field is in one direction and accelerates it when in the opposite direction.

## 5. Conductivity of Metallic Vapours in Flames.

When the vapours of certain metallic salts are introduced into a flame the electrical conductivity is greatly increased. The alkali metals of larger atomic weight produce the greatest conductivity.

It is found that the relation between the current, the potential and the distance between the electrodes, and the distribution of potential between the electrodes, for a flame made strongly conducting by an alkali salt vapour are quite similar to those for a flame free from salt. The salt increases the ionization and so the current, but does not alter the nature of the phenomena. The Hall Effect also has about the same value as in a flame free from salt. It appears that the negative ions in a salted flame are electrons, since their velocity due to a field of 1 volt per centimetre is several thousand centimetres per second, as in an unsalted flame.

Interesting effects are produced by putting a bead of salt on a platinum wire into different parts of an otherwise unsalted flame. If the bead is put in anywhere, so that the salt vapour does not come in contact with the negative electrode, then there is no appreciable effect on the current. If, however, the salt vapour is allowed to get to the negative electrode then a very large increase of the current is produced. As we have seen, most of the fall of potential between the electrodes occurs close to the negative electrode, so that we may say that nearly all the electrical resistance of the flame is in the layer close to the negative electrode. The salt vapour therefore has very little effect unless it gets into this layer at the negative electrode.

The potential drop at the negative electrode can be made small or zero by coating the electrode with lime or barium oxide. This causes the electrode to emit electrons, so that if the electrons emitted are sufficiently numerous to carry the current the potential drop disappears. The uniform gradient  $X_0$  then extends right up to the negative electrode, and the relation between the current  $C$  and potential difference  $V$  becomes roughly

$$V = ACd,$$

instead of

$$V = ACd + BC^2.$$

The gradient  $X_0$  is then nearly equal to  $V/d$ , which is much greater than before, so that the current is proportionally increased. A similar effect can be produced by putting any alkali metal salt on the negative electrode.

If the potential difference is reversed so that the coated electrode is positive, the current is then given by  $V = ACd + BC^2$ , and is the same as when both electrodes are of clean platinum. Thus with one electrode coated with lime the flame acts as a rectifier for an alternating current. When the negative electrode is coated with lime the equation  $V = ACd$  holds approximately, provided the current  $C$  is not greater than the current the electrons emitted by the lime can carry. If  $V$  is increased so that  $C$  becomes greater than this, the negative drop reappears.

When there is little or no negative drop, putting a bead of salt into

any part of the flame between the electrodes increases the current because the resistance is not then concentrated at the negative electrode. Such experiments show clearly that the salt vapour is strongly ionized in any part of the flame.

Different salts of the same alkali metal, for example  $\text{KCl}$ ,  $\text{K}_2\text{CO}_3$ , and  $\text{KNO}_3$  give nearly equal conductivities to a flame. It is therefore probable that the salts are dissociated and that the metal is present in the flame as metallic vapour. The extreme smallness of the partial pressure of the salt in the flame, which is usually of the order of  $10^{-6}$  mm., makes such a dissociation very probable.

## 6. Thermodynamical Theory.

The metallic atoms in the flame are ionized by collisions with other atoms, or by the action of light radiation, so that we get electrons and positively charged metallic atoms. The electrons and ions recombine, a state of equilibrium being established when the ionization is equal to the recombination. The condition of equilibrium can be obtained by means of the thermodynamical theory of chemical equilibrium in a mixture of gases. It is shown in the chapter on the Quantum Theory (section 7) that the entropy  $\Phi$  of one mol of a monatomic gas is given by the equation

$$\Phi = k\mathcal{N} \log \left\{ \frac{V\varepsilon^{5/2}}{\mathcal{N}\hbar^3} (2\pi mk\theta)^{3/2} \right\},$$

where  $k$  is the gas constant for one molecule,  $\mathcal{N}$  the number of molecules of any gas in one mol,  $V$  the volume of one mol of the gas,  $\varepsilon$  the base of Napierian logs,  $\hbar$  Planck's constant,  $\theta$  the absolute temperature, and  $m$  the mass of one molecule.

Consider a very large quantity of the metallic vapour and let  $p_1$  be the partial pressure of the metallic atoms in it,  $p_2$  the partial pressure of the positive ions, and  $p_3$  that of the electrons. Suppose that the mixture is in a state of equilibrium, and that one mol of the atoms dissociates into ions and electrons. Then, provided the amount of the vapour is so large that this dissociation produces no appreciable change in the partial pressure, the resulting total change of entropy must be zero, by the second law of thermodynamics. Hence

$$\Phi_3 + \Phi_2 - \Phi_1 - \frac{H}{\theta} = 0,$$

where  $\Phi_1$  is the entropy of one mol of the atoms,  $\Phi_2$  that of one mol of the ions,  $\Phi_3$  that of one mol of the electrons, and  $H$  the amount of heat energy which must be added to the vapour to keep its temperature constant during the dissociation. For  $\Phi_3 + \Phi_2 - \Phi_1$  is the increase in the entropy of the vapour, and  $H/\theta$  is the entropy lost by the surrounding bodies which supply the heat  $H$ . Now, in the metallic vapour, the ions and the electrons are all monatomic gases, so that we may substitute for  $\Phi_1$ ,  $\Phi_2$ , and  $\Phi_3$  the above expression for  $\Phi$ .

Putting  $pV = k\mathcal{N}\theta$ , it becomes

$$\Phi = k\mathcal{N} \log \left\{ \frac{k\theta\varepsilon^{5/2}}{p\hbar^3} (2\pi mk\theta)^{3/2} \right\},$$

so that we get

$$\Phi_2 - \Phi_1 = k\mathcal{N} \log \left\{ \frac{p_1}{p_2} \left( \frac{m_2}{m_1} \right)^{3/2} \right\},$$

where  $m_2$  is the mass of one ion and  $m_1$  that of one atom. But  $m_1$  and  $m_2$  are

practically equal, since the mass  $m_3$  of an electron is negligible compared with that of an atom. Hence

$$\Phi_2 - \Phi_1 = k\mathcal{N} \log \frac{p_1}{p_2}.$$

The equation  $\Phi_3 + \Phi_2 - \Phi_1 - \frac{H}{\theta} = 0$  therefore gives

$$\log \frac{p_2 p_3}{p_1} = -\frac{H}{k\mathcal{N}\theta} + \log \left\{ \frac{k\theta \varepsilon^{5/2}}{\hbar^3} (2\pi m_3 k\theta)^{3/2} \right\}.$$

Now  $H$  is equal to  $H_0 + k\mathcal{N}\theta$ , where  $H_0$  is the increase in the internal energy due to the dissociation, for  $k\mathcal{N}\theta = pV$  is the external work done. Let  $H_0 = \mathcal{N}Pe$ , where  $e$  is the charge on one electron, and  $P$  the potential difference through which a charge  $e$  must fall to acquire enough energy to dissociate one of the metallic atoms into an ion and an electron.

Then, putting  $K = \frac{p_2 p_3}{p_1}$ , we get

$$\log K = -\frac{Pe}{k\theta} + \frac{5}{2} \log \theta + \log \left\{ \frac{k^{5/2}}{\hbar^3} (2\pi \varepsilon m_3)^{3/2} \right\}.$$

All the quantities in the last term on the right-hand side of this equation are known, so that we can calculate  $K$  at any temperature  $\theta$ , provided we know  $P$  for the metal vapour.

## 7. Conductivity with Varying Amounts of Salt in Flame.

In the flame there is some ionization of the flame gases in addition to the ionization of the metal vapour. Let  $p_2'$  be the partial pressure of the positive ions formed from the flame gases, and  $p_1'$  the partial pressure of the undissociated flame molecules, and let  $K' = \frac{p_2' p_3}{p_1'}$  be the equilibrium constant for the flame molecules.

The conductivity of the flame is proportional to the number of electrons, so that we have  $p_3 = Ac$ , where  $A$  is a constant and  $c$  denotes the conductivity. Also  $p_3 = p_2 + p_2'$ , since the number of electrons must be equal to the total number of ions. Let  $p = p_1 + p_2$  be the partial pressure of the metal atoms, neutral and dissociated; also let  $p' = p_1' + p_2'$ .

We shall assume  $p_2'$  to be very small compared with  $p_1'$ , so that approximately  $p' = p_1'$ .

Now 
$$K = \frac{p_2 p_3}{p - p_2}, \text{ or } p_2 = \frac{Kp}{K + Ac}.$$

Also 
$$K' = \frac{p_2' p_3}{p'}, \text{ so that } p_2' = \frac{K'p'}{Ac}.$$

Hence the equation 
$$Ac = p_2 + p_2'$$

becomes 
$$Ac = \frac{Kp}{K + Ac} + \frac{K'p'}{Ac}.$$

When there is no metal vapour in the flame so that  $p = 0$ , let  $c = c_0$ , so that

$$Ac_0 = \frac{K'p'}{Ac_0};$$

then putting  $\frac{c}{c_0} = x$ , we get

$$A \left( x - \frac{1}{x} \right) = \frac{Kp}{c_0(K + Ac_0x)},$$

or

$$\frac{px}{x^2 - 1} = Ac_0 \left( 1 + \frac{Ac_0x}{K} \right).$$

The partial pressure  $p$  is proportional to the amount of salt in the flame, so that this equation can be tested by observing how the conductivity varies with the amount of salt in the flame. When  $x$  is large it becomes approximately  $Kp = (Ac_0x)^2$ , and the conductivity should therefore be proportional to the square root of the amount of salt present. A definite amount of any salt can be introduced into a flame by spraying a solution of the salt by means of a jet of compressed air, mixing the air and spray with coal gas and then passing it into the burner. In this way the salt is uniformly distributed through the flame, and the amount of it entering the flame is proportional to the concentration of the solution used. By finding the amount of solution used up in a known time the amount of salt entering the flame in unit time can be determined.

If  $g$  is the weight of salt per unit volume in the salt solution sprayed into the flame we may put  $p = Bg$ , where  $B$  is a constant, so that

$$\frac{gx}{x^2 - 1} = \frac{Ac_0}{B} \left( 1 + \frac{Ac_0x}{K} \right).$$

If  $x = c/c_0$  is found for a series of values of  $g$ , then on plotting  $gx/(x^2 - 1)$  against  $x$  a straight line given by  $y = b + ax$  should be obtained, where  $y = \frac{gx}{x^2 - 1}$ ,  $b = \frac{Ac_0}{B}$ , and  $a = \frac{A^2c_0^2}{BK}$ . Hence  $K = B \frac{b^2}{a}$ . The conductivity  $c$  of the flame is proportional to the ratio of the current to the uniform potential gradient between the electrodes, so that  $x = c/c_0$  can be easily determined.

The following table gives the results of a series of measurements of  $x$  for a flame into which solutions of caesium chloride were sprayed.

Grammes CsCl per Litre in Solution ( $g$ ).	$x$ .	$\frac{10^4gx}{x^2 - 1}$ .	$10 + x$ .
0	1	—	—
0.0032	2.88	12.6	12.83
0.008	5.72	14.5	15.72
0.016	8.9	18.2	18.9
0.032	13.5	23.9	23.5
0.08	22.7	36.0	32.7
0.16	32.8	49.0	42.8
0.8	85.2	94.0	95.2
8.0	282.0	284.0	292.0
80.0	883.0	906.0	893.0

It appears that  $\frac{10^4gx}{x^2 - 1}$  is nearly equal to  $10 + x$ , so that the conductivity

of the flame varies with the amount of metal vapour in it, approximately in accordance with the ionic theory. Similar results have been obtained with other alkali metals.

The equation  $p_2 = \frac{Kp}{K + Ac}$  gives for the fraction of the metal atoms ionized  $\frac{p_2}{p} = \frac{K}{K + Ac}$ , which is equal to  $\frac{b}{b + ax}$ . In the case of caesium therefore in the particular flame used the fraction of the caesium atoms ionized was  $\frac{10}{10 + x}$ . With an indefinitely small amount of caesium  $x = 1$ , so that the fraction ionized is 10/11. The ionization does not become complete because there are some electrons in the flame when no caesium is present.

If the number of metal atoms in unit volume of the flame is  $n$  and the corresponding conductivity  $c$ , the number of electrons in unit volume is  $n_3 = nbx/g$ , for we have  $p_3 = Ac_0x$  and  $p = Bg$ , and therefore

$$\frac{p_3}{p} = \frac{n_3}{n} = \frac{Ac_0x}{Bg},$$

and  $b = Ac_0/B$ . Thus the number  $n_3$  of electrons in unit volume of the flame can be calculated from the ratio  $n/g$ . To find  $n/g$  it is necessary to find the amount of solution entering the flame, and the volume of the flame gases with which it is mixed. The volume of the flame gases can be found from the horizontal cross-section of the flame and the upward velocity of the flame gases. In this way  $n/g$  can be estimated and so  $n_3$  found. The current density in the flame in the uniform potential gradient is  $n_3ek_2X_0$ , so that when  $n_3$  is known  $k_2$ , the mobility of the electrons, can be obtained. In this way it has been found that  $k_2$  is about 4000 cm. per second for 1 volt per centimetre.

The equilibrium constant  $K$  can be calculated by means of  $K = Bb^2/a$ . We have  $p = Bg = nb\theta$ , where  $n$  is the number of metal atoms per unit volume in the flame,  $b$  the gas constant for one molecule, and  $\theta$  the absolute temperature. This gives  $B$  when  $n$  is known, and  $K$  can then be calculated. It is found that the values of  $K$  found in this way for the alkali metals are nearly equal to those given by the thermodynamical theory of the equilibrium between the atoms, ions, and electrons. This shows that the quantum theory of the entropy of electron gas is approximately correct.

## 8. Conductivity for Alternating Currents.

The conductivity of flames for rapidly alternating currents has been investigated with interesting results. Suppose we have two plane parallel electrodes in a Bunsen flame at a distance  $d$  apart, and that an alternating potential difference is maintained between them. Let the electrodes be symmetrically placed in the flame and red hot.

The mobility of the positive ions in the flame is very small compared with that of the electrons, and their mass is enormously greater. Clearly, then, in a rapidly alternating electric field the amplitude of vibration of the ions will be very small compared with that of the electrons. We shall therefore assume that the positive ions do not move at all and that the current is all carried by the electrons. When there is no electric field, let there be  $n$  positive ions per unit volume throughout the space between the electrodes, and an equal number of

electrons. When the alternating field is applied let the electrons vibrate with an amplitude  $A$ . All the electrons within a distance  $A$  of the electrodes will therefore strike the electrodes and be removed from the flame. We suppose  $2A$  less than  $d$ , so that a layer of electrons of thickness  $d - 2A$  remains in the flame and oscillates between the electrodes.

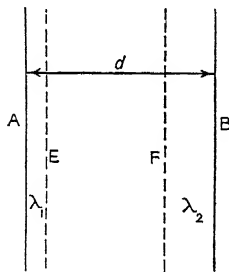


Fig. 2

In the layer of electrons the density of charge is zero, but outside the layer it is  $ne$  where  $e$  is the charge on one positive ion.

In fig. 2 A and B are the electrodes, and the layer of electrons is between E and F. Let the layer near A in which there are no electrons be of thickness  $\lambda_1$ , and that near B of thickness  $\lambda_2$ . Then  $\lambda_1 + \lambda_2 = 2A$ , and if  $y$  denotes the displacement of the layer of electrons from its mean position then

$$\lambda_1 - \lambda_2 = 2y.$$

Let  $V$  be the potential difference between the electrode A and a point at a distance  $x$  from it. From  $x = 0$  to  $x = \lambda_1$  and from  $x = d - \lambda_2$  to  $x = d$ , we have  $\frac{\partial^2 V}{\partial x^2} = -4\pi\rho$  where  $\rho = ne$ . Also from  $x = \lambda_1$  to  $x = d - \lambda_2$  we have  $\frac{\partial^2 V}{\partial x^2} = 0$ . By integrating these equations it is easy to calculate  $V$  at  $x = d$ . The integration constants are found by making  $V$  and  $\frac{\partial V}{\partial x}$  continuous at  $x = \lambda_1$  and  $x = d - \lambda_2$ . If  $X$  denotes the field strength in the space between E and F, we find for the potential difference between the electrodes

$$V = 8\pi\rho Ay - Xd.$$

The term  $Xd$  will probably be small unless  $\rho$  is very small, so that approximately when  $\rho$  is large  $V = 8\pi\rho Ay$ .

The equation of motion of the electrons is

$$m \frac{d^2 y}{dt^2} = -Xe - \mu \frac{dy}{dt},$$

where  $m$  is the mass and  $-e$  the charge of one electron, and  $\mu$  is the average viscous resistance to the motion for unit velocity. We shall suppose that  $\mu$  is small, so that approximately

$$m \frac{d^2 y}{dt^2} = -Xe.$$

If then  $X = X_0 e^{ipt}$ , and  $y = A e^{ipt}$ , then  $A = eX_0/mp^2$ . Putting  $V = V_0 e^{ipt}$ , and substituting in  $V = 8\pi\rho Ay$ , we get

$$A = \sqrt{\frac{V_0}{8\pi\rho}}.$$

The current density  $I$  is given approximately by

$$I = -\rho \frac{dy}{dt} = -\rho \sqrt{\frac{V_0}{8\pi\rho}} ip e^{ipt}.$$

This shows that the electrodes in the flame behave like a condenser. For the charge on a condenser of capacity  $C$  is  $Q = VC = CV_0 e^{ipt}$ , and the current charging it is  $-\frac{dQ}{dt} = -CV_0 ip e^{ipt}$ . Comparing this with the expression for  $I$  we see that the electrodes in the flame have an apparent capacity  $C$  per unit area given by

$$C = \sqrt{\frac{\rho}{8\pi V_0}}.$$

The apparent capacity of the electrodes in the flame can be measured by means of a high-frequency Wheatstone bridge in which the capacity of the electrodes is balanced by three air condensers. In this way it has been found that  $C$  is nearly inversely as  $\sqrt{V_0}$  and directly as  $\sqrt{\rho}$ , in accordance with the theory. If the viscous resistance to the motion of the electrons is not neglected, then the theory indicates that the electrodes should behave like a condenser shunted by a high resistance. By determining the apparent resistance it is possible to estimate the viscous resistance to the motion of the electrons and so get an estimate of their mobility. The results obtained in this way are of the same order as those given by the other methods.

### 9. Mobility of Positive Ions in Flames.

The mobility of the positive ions of alkali metals in flames has been estimated by finding the strength of the electric field required to make them move down the flame against the upward stream of gases.

In fig. 3 FF is a Bunsen flame. A and B are two electrodes consisting of fine platinum wire gratings with wires about 0.5 cm. apart which are supported one above the other in the flame. A bead of salt on a wire D can be introduced into the flame just below the upper electrode. The upper electrode is charged positively and the current from it to a wire C, the end of which is in the plane of B at the axis of the flame, is measured with a galvanometer. The wire C is connected to one terminal of the galvanometer and the other terminal is connected to the electrode B, which is connected to the earth. The current is measured with and without the salt bead in the flame. It is found

that when the potential difference between A and B exceeds a certain value, the current down the axis of the flame to C is increased by putting in the bead. This is supposed to indicate the point at which the ions from the salt move down. When care is taken to prevent salt vapour from the bead getting into the lower parts of the flame, it is found that the electric field required to make the ions move down is about 200 volts per centimetre. The upward velocity of the flame gases is about 200 to 300 cm. per second, so that the mobility of the ions is about 1 cm. per second for 1 volt per centimetre. The mobility of the electrons

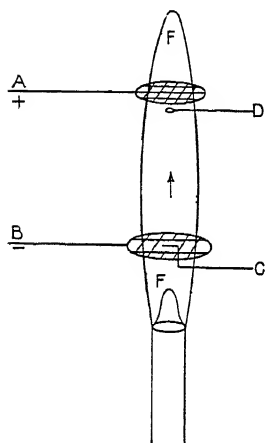


Fig. 3

being about 4000 cm. per second for 1 volt per centimetre, it follows that practically all the current is carried by the electrons, except close to the negative electrode. It is found that the mobility of the positive ions is practically the same for lithium, sodium, potassium, rubidium, and caesium ions.

The mobility of an electron is nearly equal to  $e\lambda/mV$ , where  $e$  is the charge and  $m$  the mass of an electron,  $\lambda$  the mean free path, and  $V$  the average velocity of agitation. This expression was obtained by assuming that after a collision of an electron with a molecule the velocity of the electron was equally likely to be in any direction. This assumption cannot be made in the case of an ion, because the mass of an ion is comparable with that of a molecule.

If the velocity of an ion is given by  $v = kX$ , where  $X$  is the field, the force on an ion moving with unit velocity through the gas must be  $Xe/v = e/k$ . If an ion of mass  $M$  is projected into the gas with initial velocity  $V_0$ , the average distance it will travel in its original direction may be calculated as follows.

Let a large number  $N$  of ions be projected into the gas with average initial velocity  $V_0$ . Then

$$\frac{d}{dt}(NMV) = -\frac{e}{k}NV,$$

where  $V$  is the average velocity of the  $N$  ions in the original direction. This gives

$$V = V_0 e^{-\frac{e}{kM}t}.$$

The average distance  $L$  which they go in the original direction is

$$L = \int_0^\infty V dt = V_0 \int_0^\infty e^{-et/kM} dt = \frac{V_0 kM}{e}.$$

Thus  $k = \frac{eL}{MV_0}$ , so that if  $V_0$  is equal to the average velocity of agitation of the ions, which we denote by  $V$ , then

$$k = \frac{eL}{MV}.$$

In the case of an electron  $L = \lambda$ , since the electrons lose all their original velocity on the average at the first collision. The positive ions of the different alkali metals in a flame all have about equal mobilities, so that it appears that  $L$  is proportional to  $MV$ , or to  $\sqrt{M}$ , since  $V$  is inversely as  $\sqrt{M}$ . The mobility of the positive ions in flames seems to be about the same as that of the positive ions produced by X-rays in air at atmospheric pressure and at the ordinary temperature.

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## CHAPTER XVI

# The Positive Column and Negative Glow

### 1. The Positive Column, Negative Glow, Crookes Dark Space, and Faraday Dark Space.

When an electric discharge is passed between electrodes in a gas at a pressure of the order of 1 mm. of mercury, a luminous column extending some distance from the positive electrode or anode is usually obtained which is called the positive column. Near the negative electrode or cathode there is a luminous layer separated from the cathode by a non-luminous region which is called the Crookes dark space. The luminous layer is sharply defined on the side near the cathode, but fades gradually on the other side with increasing distance from the cathode. This luminous layer is called the negative glow.

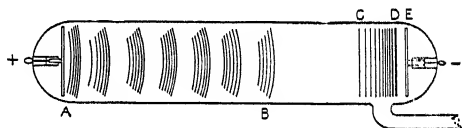


Fig. 1

Between the negative glow and the positive column there is a non-luminous region called the Faraday dark space. The colour of the light emitted by the negative glow is often quite different from that emitted by the positive column.

When the distance between the electrodes is increased, keeping the gas pressure and the current constant, the distances of the negative glow and of the end of the positive column from the cathode remain unchanged, so that the length of the positive column increases by an amount equal to the increase in the distance between the electrodes.

The luminosity of the positive column is sometimes uniform along its whole length, but frequently varies periodically so that it consists of a series of equally spaced bright layers or striæ separated by more or less dark intervals. Large currents and low pressures favour the appearance of striæ. Fig. 1 shows such a discharge in nitrogen gas. The discharge is produced in a glass tube about 2 cm. in diameter and 20 cm. long, having aluminium disc electrodes sealed in at each end as shown. The positive column, consisting of 8 striæ, extends from A to

B. The Faraday dark space is between B and C, the negative glow between C and D, and the Crookes dark space between D and E. The pressure is about  $\frac{1}{4}$  mm. and the current about 5 milliamperes in such a discharge as this.

As the pressure in such a discharge tube is reduced, the positive column gets shorter and finally disappears except for a layer of luminosity on the anode. The negative glow and Crookes dark space get longer, and eventually the Crookes dark space extends to the anode when the pressure is about 0.003 mm. The potential difference required to maintain the discharge depends on the gas pressure. With a discharge tube 3 cm. in diameter having electrodes 11.5 cm. apart and a constant current of 10 milliamperes Townsend obtained the following results:

Pressure in millimetres of mercury .. .. .	}	4.0	2.84	1.65	1.04	0.66	0.4	0.29	0.24	0.13
Potential difference in volts		650	620	500	470	490	530	590	630	800

As the pressure is reduced the potential difference falls to a minimum and then rises again. At very low pressures the potential difference required to produce a discharge becomes very large, and it is possible to obtain so good a vacuum that two hundred thousand volts will not produce any discharge.

## 2. Potential Differences in the Tube.

The potential at any point in a discharge tube may be estimated by means of a small insulated electrode of fine wire projecting into the tube. The wire is usually enclosed in a glass tube except near its end. The potential difference between the wire and one of the electrodes can be measured with an electrostatic voltmeter. Such an insulated

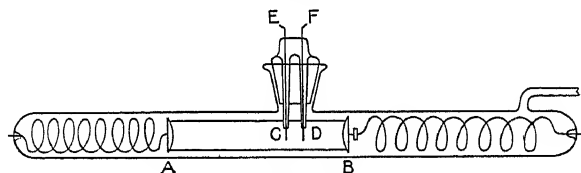


Fig. 2

wire takes up a definite potential which is probably not much different from the potential of the discharge close to it. The current in the discharge is carried by electrons moving towards the anode and positive ions moving towards the cathode. The electrons have much higher velocities than the ions. The wire electrode takes up a potential such that the negative charge it receives by absorbing electrons is equal to the positive charge it receives from the positive ions. It therefore probably takes up a potential somewhat below that of the gas near it.

so that it repels electrons and attracts ions, for owing to the higher velocity of the electrons more electrons move towards the wire than ions.

The strength of the electric field along the discharge can be found by means of two similar small electrodes a few millimetres apart. The potential difference between them can be measured with an insulated

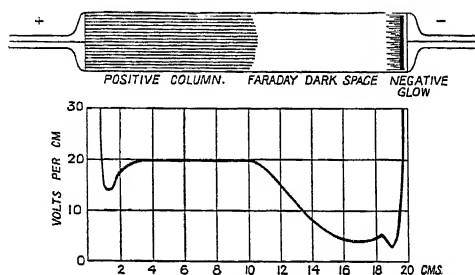


Fig. 3

quadrant electrometer, and the field strength is equal to the potential difference divided by the distance between the electrodes.

A discharge tube used by the writer for such measurements is shown in fig. 2. A and B are two aluminium disc electrodes kept at a constant distance apart by three thin glass rods, and connected to wires sealed through the ends of the tube by spiral springs of fine wire. A small piece of iron enables the electrodes to be moved along the tube by means of a magnet. C and D are two fine wire electrodes enclosed in small glass tubes except close to their ends and sealed into a glass stopper as shown. By turning the stopper the distance between the electrodes measured along the axis of the tube can be varied. The

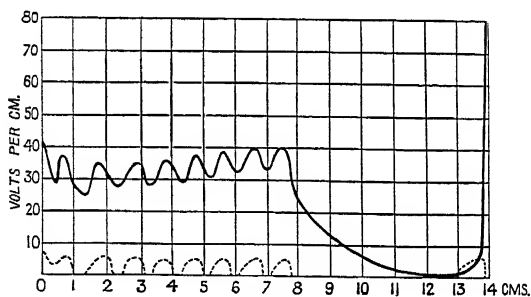


Fig. 4

electrodes C and D were connected to an insulated quadrant electrometer, and a discharge was passed between A and B from a battery of a large number of small cells.

The distribution of the electric field along the discharge is shown in figs. 3

and 4, which represent results obtained by Graham and by the writer respectively.

In fig. 3 the field strength is constant along the uniform positive column. It rises rapidly near both electrodes and is small in the negative glow and Faraday dark space.

In fig. 4 the dotted curve shows approximately the distribution of luminosity. In this case the field strength rises and falls in the positive

column, the maxima being in the striæ and the minima between them.

In discharges at rather low pressure, about 0.1 mm., when the Faraday dark space extends almost to the anode, the field strength close to the anode becomes very small or even negative.

### 3. The Cathode Fall of Potential.

There is always a considerable difference of potential between the negative glow and the cathode. This is called the cathode fall of potential. There is also a smaller fall of potential very close to the surface of the anode. The electric field is therefore very large and positive close to the surface of the anode but is very small or even negative 1 or 2 mm. from it. The variation of the field near the anode is similar to that near the cathode but is spread over a much smaller space. At the cathode positive ions strike the metal surface and liberate electrons, whereas at the anode electrons strike the metal surface and may liberate some positive ions from it.

The cathode fall of potential can be determined by measuring the potential difference between the cathode and a small wire electrode in the negative glow by means of an electrostatic voltmeter. The negative glow does not cover the whole surface of the cathode when the current is not too large. As the current is increased, the area of the cathode covered by the negative glow is proportional to the current, so that the current density in the glow remains constant. When the glow covers the whole cathode it becomes brighter as the current is increased. The cathode fall of potential is independent of the current and of the gas pressure so long as the cathode is only partially covered by the glow, but increases with the current when the cathode is covered. The cathode fall of potential when the cathode is only partly covered is called the normal cathode fall of potential.

The following table gives some values of the normal cathode fall of potential for cathodes made of different metals.

Gas.	Platinum.	Aluminium.	Potassium.
Oxygen ..	369 volts	—	—
Hydrogen ..	295 "	190 volts	172 volts
Nitrogen ..	232 "	224 "	170 "
Helium ..	226 "	—	60 "
Argon ..	167 "	—	—

The normal cathode fall of potential is nearly equal to the minimum sparking potential, that is, the sparking potential between parallel plate electrodes when the gas pressure is adjusted so that the sparking potential is as small as possible. The large influence of the nature of

the cathode on the normal cathode fall of potential is probably due to the emission of electrons by the cathode when bombarded by positive ions.

Aston and Watson made a number of measurements of the potential difference required to maintain a discharge between parallel plate electrodes. In their experiments the negative glow extended to the anode so that there was no positive column, and the potential difference between the electrodes was practically equal to the cathode fall of potential. The cathode was entirely covered by the glow, so that the cathode fall of potential was greater than the normal value.

It was found that the potential difference  $V$  was given by the following equation

$$V = E + \frac{F\sqrt{C}}{p},$$

where  $E$  and  $F$  are constants,  $C$  is the current density, and  $p$  the gas pressure.

Aston and Watson also measured the length  $D$  of the Crookes dark space and found that

$$D = \frac{A}{p} + \frac{B}{\sqrt{C}},$$

where  $A$  and  $B$  are constants. The values of the constants  $E$ ,  $F$ ,  $A$ , and  $B$  depend on the nature of the gas and of the cathode. With  $D$  in centimetres,  $p$  in  $\frac{1}{100}$  mm. of Hg,  $C$  in  $10^{-4}$  ampere per square centimetre, the following table gives some values of these constants.

Cathode.	Oxygen.				Hydrogen.			
	$A$ .	$B$ .	$E$ .	$F \times 10^{-2}$ .	$A$ .	$B$ .	$E$ .	$F \times 10^{-2}$ .
Aluminium	5.7	0.43	310	17.5	23	0.41	170	66
Copper ..	8.9	0.40	340	28.5	47	0.45	300	130
Platinum..	8.8	0.40	335	30.0	45	0.42	270	120

It appears that, for a given gas and cathode,  $V$  and the product  $pD$  are functions of  $\sqrt{C}/p$ . If we eliminate  $\sqrt{C}/p$  from the equations giving  $V$  and  $D$  we get

$$V = E + \frac{FB}{pD - A},$$

so that  $V$  depends on  $pD$  only in any given case. The product  $pD$  is proportional to the amount of gas per unit area in the Crookes dark space, and  $V$  is nearly equal to the fall of potential across the Crookes

dark space, since the field strength in the negative glow is quite small. It is also found that the sparking potential between parallel plates is a function of the amount of gas per unit area between the plates.

The electric field strength in the Crookes dark space has been investigated by several physicists by observing the potentials taken up by a wire electrode at different distances from the cathode. This method is unreliable when the wire is close to the cathode, because the wire stops the stream of positive ions moving towards the cathode and so prevents the emission of electrons by the cathode immediately behind the wire. The wire therefore becomes positively charged and so does not take up the potential of the gas.

Aston determined the electric field in the Crookes dark space by measuring the deflection of a narrow beam of cathode rays sent across the dark space parallel to the surface of the cathode. In this way he found that the field strength is nearly proportional to the distance from the negative glow, so that  $F = Ax$ , where  $F$  is the field strength,  $x$  the distance from the glow, and  $A$  a constant. The cathode fall of potential  $V$  is therefore given by

$$V = \int_0^D Ax dx = \frac{AD^2}{2}.$$

Aston determined  $A$  by measuring  $F$  at different distances from the negative glow and found that  $\frac{1}{2}AD^2$  was equal to the cathode fall of potential determined by measuring the potential difference between the negative glow and the cathode.

#### 4. Theory of the Cathode Fall of Potential.

An approximate theory of the cathode fall of potential can be worked out if we assume that the gas in the Crookes dark space is ionized by collisions with electrons and with positive ions, and that electrons are liberated by collisions of positive ions with the cathode. We also assume that recombination of the ions and electrons can be neglected, and that the number of ions and electrons lost by diffusion to the walls of the discharge tube is inappreciable. The condition which must be satisfied in order that a steady current may be maintained through a gas between parallel electrodes with these assumptions, when no electrons are set free at the cathode, was worked out by Townsend. A slight modification of Townsend's theory gives the condition when electrons are supposed set free at the cathode.

Let  $n_1$  be the number of positive ions per cubic centimetre at a point, and  $v_1$  their velocity; and let  $n_2$  and  $v_2$  be the corresponding quantities for the negative electrons. Let  $\alpha$  denote the number of molecules ionized by a negative electron in moving 1 cm., and let  $\beta$  be the corresponding number for a positive ion. The electric intensity  $X$  varies with the distance  $x$  from the cathode, so that  $\alpha$ ,  $\beta$ ,  $v_1$ , and  $v_2$  are not constant. In the steady state we have

$$-\frac{d}{dx}(n_2v_2) + \alpha n_2v_2 + \beta n_1v_1 = 0,$$

and

$$\frac{d}{dx}(n_1v_1) + \alpha n_2v_2 + \beta n_1v_1 = 0.$$

Also, the current density  $C$  is equal to  $e(n_1v_1 + n_2v_2)$  and is constant. Substituting  $C/e - n_1v_1$  for  $n_2v_2$ , we get

$$\frac{d}{dx}(n_1v_1) - (\alpha - \beta)n_1v_1 = -\alpha C/e.$$

Let

$$e \int_0^x (\alpha - \beta) dx = Z,$$

so that

$$\frac{d}{dx}(n_1v_1 Z^{-1}) = -\frac{\alpha CZ^{-1}}{e}.$$

Then

$$n_1v_1 = BZ - \frac{C}{e} Z \int_0^x \alpha Z^{-1} dx,$$

where  $B$  is constant.

The number of positive ions striking the cathode per square centimetre per second is equal to  $n_1v_1$  at  $x=0$ . Let  $\gamma n_1v_1$  be the number of electrons set free by the impacts of these positive ions on the cathode. Then at  $x=0$  we have  $n_2v_2 = \gamma n_1v_1$ , or

$$n_1v_1 = \frac{C}{(1 + \gamma)e}.$$

This condition gives

$$B = \frac{C}{(1 + \gamma)e}.$$

Let the distance between the electrodes be  $S$ , so that at  $x=S$ ,  $n_1v_1=0$ . This gives

$$0 = \frac{CZ}{e(1 + \gamma)} - \frac{CZ}{e} \int_0^S \alpha Z^{-1} dx,$$

or

$$\int_0^S \alpha Z^{-1} dx = \frac{1}{1 + \gamma},$$

which is the condition required. If  $\gamma=0$ , this reduces to the condition given by Townsend.

In a discharge at moderately low pressure, when a Crookes dark space exists, the electric intensity is very small in the negative glow, so that if the positive electrode is anywhere in the negative glow then

$$\int_0^S \alpha Z^{-1} dx = \int_0^D \alpha Z^{-1} dx,$$

where  $D$  is the length of the Crookes dark space, because  $\alpha$  and  $\beta$  are both zero when  $X$  is very small.

Let  $\alpha = \alpha_1 p$ ,  $\beta = \beta_1 p$ ,  $x = y/p$ , and  $Z_1 = e \int_0^y (\alpha_1 - \beta_1) dy$ , so that

$$\int_0^D \alpha Z^{-1} dx = \int_0^{pD} \alpha_1 Z_1^{-1} dy = \frac{1}{1 + \gamma}.$$

Also let the field strength  $X = pY$ . The equation  $X = A(D - x)$  becomes  $Y = \frac{A}{p^2}(pD - y)$ , and, putting  $A_1 = A/p^2$ , we get  $Y = A_1(pD - y)$ . The equation

$V = \int_0^D X dx$  then becomes

$$V = \int_0^{pD} Y dy = A_1 \frac{(pD)^2}{2}.$$

For the normal cathode fall of potential in hydrogen, Skinner found  $V = 197$  volts, and  $pD = 1.10$ . In this case, therefore,  $A_1 = \frac{2 \times 197}{(1.10)^2} = 325$ , whence  $Y = 325(pD - y)$ .

Now Townsend has determined  $\alpha$  and  $\beta$  as functions of  $Y = X/p$ , so that by using his values of these quantities we can compute the value of  $\int_0^{pD} \alpha_1 Z_1^{-1} dy$ . It is found in this way that this integral is nearly equal to unity, and  $\gamma$  must therefore be very small.

The equation  $n_1 v_1 = BZ - \frac{C}{e} Z \int_0^x \alpha Z^{-1} dx$ , with  $B = \frac{C}{(1 + \gamma)e}$ , gives

$$n_1 v_1 = \frac{C}{e} Z \left( \frac{1}{1 + \gamma} - \int_0^x \alpha Z^{-1} dx \right),$$

where  $x$  is the distance from the cathode. Also  $n_2 v_2 = \frac{C}{e} - n_1 v_1$ . Hence, substituting for  $n_1$  and  $n_2$  in the equation

$$\frac{dX}{dx} = 4\pi e(n_1 - n_2),$$

we get, on putting  $v_1 = k_1 X$  and  $v_2 = k_2 X$ ,

$$\frac{dX^2}{dx} = -8\pi C \left\{ \frac{1}{k_2} - \left( \frac{1}{k_1} + \frac{1}{k_2} \right) Z \left( \frac{1}{1 + \gamma} - \int_0^x \alpha Z^{-1} dx \right) \right\}.$$

Let  $X = pY$ ,  $px = y$ ,  $k_2 = K_2/p$ ,  $k_1 = K_1/p$ ,  $\alpha = \alpha_1 p$ ,  $\beta = \beta_1 p$ ; then

$$\frac{dY^2}{dy} = -\frac{8\pi C}{p^2} \left\{ \frac{1}{K_2} - \left( \frac{1}{K_1} + \frac{1}{K_2} \right) Z_1 \left( \frac{1}{1 + \gamma} - \int_0^y \alpha_1 Z_1^{-1} dy \right) \right\}.$$

where  $Z_1 = e^{\int_0^y (\alpha_1 - \beta_1) dy}$ , as before.

Now  $K_2$ ,  $K_1$ ,  $\alpha_1$ , and  $\beta_1$  are all functions of  $Y$  only, for Townsend has proved experimentally that  $\alpha/p$ ,  $\beta/p$ ,  $v_1$ , and  $v_2$  are all functions of  $X/p$ . Also  $\gamma$  is presumably a function of  $X/p$  only, since it must depend on the velocity of the positive ions at  $x = 0$ . It appears therefore that the above equation is a relation between the three quantities  $Y$ ,  $y$ , and  $C/p^2$ . Hence we may write

$$Y = \varphi(y, C/p^2),$$

where  $\varphi$  denotes some function of  $C/p^2$  and  $y$  only.

Now the cathode fall of potential  $V$  is equal to  $\int_0^p X dx$ , and therefore

$$V = \int_0^{pD} Y dy.$$

For a given value of  $C/p^2$ ,  $Y$  is a function of  $y$  only, so that  $pD$  is a function of  $C/p^2$  only, for  $pD$  is the value of  $y$  at which  $Y$  becomes very small. Hence it follows that  $V$  is a function of  $C/p^2$  only. If  $V$  has a minimum value when  $C/p^2$  is varied, this minimum value will be independent of  $p$ , and the corresponding values of  $C$  will be proportional to  $p^2$ , and the corresponding values of  $D$  will be inversely as  $p$ .

## 5. Comparison with Experiment.

These results deduced from the theory are precisely those which follow from the experiments of Aston and Skinner. Thus Aston found  $pD$  and  $V$  both to

depend on  $C/p^2$ , and Skinner found the current density to be proportional to  $p^2$ , and  $D$  to be inversely as  $p$  for the normal cathode fall, which must be a minimum value of  $V$ .

At  $y = 0$ , the equation for  $\frac{dY^2}{dy}$  gives approximately

$$\frac{dY^2}{dy} = -\frac{8\pi C}{K_1 p^2},$$

since  $K_2$  is very large compared with  $K_1$ .

If now we assume that  $Y = A_1(pD - y)$ , we get

$$\frac{dY^2}{dy} = -2A_1^2(pD - y).$$

At  $y = 0$ , this gives

$$\frac{dY^2}{dy} = -2A_1^2 pD.$$

Also  $V = \int_0^{pD} Y dy$ , so that  $A_1 = 2V/p^2 D^2$ ,

and therefore

$$K_1 = \frac{\pi C (pD)^3}{V^2 p^2}.$$

The following table gives the values of  $pD$ ,  $C/p^2$ , and  $V$  found by Skinner in hydrogen, and the calculated values of  $K_1$ .  $C$  is in milliamperes per square centimetre,  $D$  in centimetres, and  $p$  in millimetres of mercury.

$C/p^2$ .	$pD$ .	$V$ .	$K_1$ .
0.0742	1.100	197	$7.2 \times 10^3$
0.1484	0.844	204	6.1 „
0.2968	0.624	227	3.9 „
		Mean	$5.7 \times 10^3$

The values of  $K_1$  are for 1 volt per centimetre at 1 mm. pressure.

If we assume that the velocity of the positive ions is inversely as the pressure, this gives, for the velocity due to 1 volt per centimetre at 760 mm., 7.5 cm. per second. This is nearly equal to the velocity of positive ions in hydrogen at 760 mm. as found by Zeleny, Langevin, and others.

It appears that the results obtained in the Crookes dark space are consistent with the theory that the ionization is due to collisions, and that recombination and diffusion can be neglected.

The ionization due to the positive ions  $\beta$  is small compared with that due to the electrons, so that if we put  $\beta = 0$ , the value of the integral  $\int_0^{pD} \alpha_1 Z_1^{-1} dy$  is not much affected, but must then be less than unity so that  $\gamma$  may not be zero. It is probable that  $\gamma$  is not exactly zero, and it is difficult to distinguish between ionization by collisions of the ions with the gas molecules, and the liberation of electrons from the cathode by the impact of positive ions. It is quite possible that  $\beta$  is really zero, and that discharges in gases are maintained by the liberation of electrons from the cathode and not by ionization by collisions of positive ions and gas molecules.

## 6. The Positive Column.

The electric intensity  $X$  in the uniform positive column depends on the gas pressure  $p$ , the current density  $C$ , and the diameter of the discharge tube. In wide tubes, the positive column may not fill the whole cross-section of the tube, and then has a certain cross-section which depends on the pressure and total current. This indicates that as the current density increases the electric intensity diminishes to a minimum value, and then increases as the current density is further increased. The cross-section of the positive column when it does not fill the tube increases with the total current, and the current density in it probably does not vary much as the current is increased.

The electric intensity  $X$  is nearly proportional to the square root of the pressure at pressures up to 2 or 3 mm., but at higher pressures is nearly a linear function of the pressure. For example, in air from  $p = 0.2$  mm. to  $p = 2.82$  mm.,  $X = 35\sqrt{p}$ , and in hydrogen from  $p = 0.25$  mm. to  $p = 1.36$  mm.,  $X = 28\sqrt{p}$ .

When the current density is very small  $X$  increases rapidly with the current, but soon attains a nearly constant value as the current is increased. With larger current densities above about 1 milliampere per square centimetre the electric intensity diminishes slightly as the current is increased.

In a uniform positive column the equation  $C = e(n_1v_1 + n_2v_2)$  becomes  $C = ne(v_1 + v_2)$ , because  $\frac{dX}{dx} = 4\pi e(n_1 - n_2) = 0$ , so that  $n_1 = n_2 = n$ . The velocity of the electrons  $v_2$  due to the electric field  $X$  is much greater than  $v_1$ , so that approximately  $C = nev_2$ . The velocity  $v_2$  has been determined by Townsend, and is a function of  $Y = X/p$ . Since  $X$  is nearly independent of  $C$ , except when  $C$  is very small, it follows that  $n$  must be proportional to  $C$ . The rate of production of electrons and ions in a uniform column must be equal to the rate at which they disappear by recombination and diffusion to the walls of the tube. The rate of production may be assumed to be proportional to the electrical energy dissipated per cubic centimetre, or to  $CX$ . The electric field does work on the electrons, giving them kinetic energy. When the velocity of an electron is small it loses very little energy by collisions, but when the velocity exceeds a critical value collisions result in ionization or excitation of the molecule with loss of the kinetic energy of the electron. Since  $C$  is proportional to  $n$ , and  $X$  is nearly constant, it follows that the rate of production of ions and electrons, and therefore also the rate of disappearance, must be nearly proportional to  $n$ . Recombination is proportional to  $n^2$ , so that it appears that the loss by diffusion to the walls of the tube must be large compared with the loss by recombination. We have therefore approximately  $\frac{dn}{dt} =$

$ACX - Bn$ , where  $ACX$  represents the rate of production of ions, and  $Bn$  the rate of loss. In the uniform column in a steady state  $\frac{dn}{dt} = 0$ , so that  $X = \frac{Bn}{AC} = \frac{Bn}{Anev_2} = \frac{B}{Aev_2}$ . If we take  $v_2 = K \frac{X}{p}$ , this gives

$$X = \frac{Bp}{AeKX},$$

or 
$$X^2 = \frac{B}{AeK} p.$$

According to this, when  $X$  varies as  $\sqrt{p}$ ,  $B/AeK$  must be independent of the pressure  $p$ .

At low pressures with large currents the positive column becomes striated. No satisfactory theory of the striations has been worked out, but in some cases it appears that the potential difference between successive striations is about equal to the ionization potential of the gas, that is, the potential through which an electron must fall to acquire enough energy to ionize a gas molecule by colliding with it.

In a transverse magnetic field the positive column is deflected sideways like a flexible conductor carrying a current. If the positive column is in a tube of circular cross-section which it fills completely, a uniform transverse magnetic field causes a concentration of the luminosity on one side of the tube, so that the luminosity is greatest on one side and fades away gradually towards the other side. A transverse electric field or Hall Effect is produced by the magnetic field in the plane perpendicular to the magnetic field. This Hall Effect can be examined with the apparatus of fig. 2 (p. 269). The glass stopper is turned until the two small electrodes C and D are at the same potential so that they are in a plane perpendicular to the current. When a magnetic field is then produced parallel to the electrodes C and D a potential difference is produced between them which can be measured with an insulated quadrant electrometer. If  $Z$  denotes the transverse electric field,  $H$  the magnetic field, and  $p$  the gas pressure, then in air between  $p = 0.26$  mm. and  $2.9$  mm. the writer found  $Z = 0.0248 H/p$ . In hydrogen,  $Z = 0.0205 H/p$ , and in oxygen,  $Z = 0.0379 H/p$ .

The current is carried almost entirely by the negative electrons, so that the force on the electrons due to the magnetic field must be equal and opposite to that due to the transverse electric field  $Z$ . Hence, if  $v$  is the velocity of the electrons along the tube, we have

$$Hev = Ze,$$

or 
$$v = \frac{Z}{H}.$$

Since  $Z = A \frac{H}{p}$ , where  $A$  is a constant, we get  $v = A/p$ , so that the velocity of the electrons along the positive column is inversely as the gas pressure. For air,  $A = 0.025$  with  $Z$  in volts per centimetre, so that with  $Z$  in electromagnetic units  $A = 25 \times 10^5$ , and  $v = \frac{25 \times 10^5}{p}$  cm. per second. At 1 mm. pressure in air  $X = 35$  volts per centimetre, and  $X/p = 35$ . For  $X/p = 35$ , Townsend found for the velocity of electrons in air about  $120 \times 10^5$  cm. per second, which is about five times that just deduced from the Hall Effect. This probably indicates that the theory of the Hall Effect just given needs modification. It is found that the Hall Effect in discharge tubes varies along the discharge in a similar way to the electric intensity. It is large in the Crookes dark space, and small in the negative glow and Faraday dark space.

The electrical conductivity of the discharge can be measured by measuring the current due to a small potential difference between two small electrodes a few millimetres apart. In this way it is found that the conductivity is small in the Crookes dark space and very large relatively in the negative glow. It is greater in the positive column than in the Faraday dark space. Roughly speaking, the conductivity is inversely as the electric intensity along the discharge, as we should expect.

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## CHAPTER XVII

# Atmospheric Electricity

### 1. Vertical Field in the Atmosphere. Conductivity.

Besides thunderstorms other less noticeable electrical phenomena are observed in the open air. There is usually a vertical electric field of the order of 150 volts per metre. The difference of potential between the earth and a point in the air above it may be found by means of an insulated conductor provided with some device to bring it to the same potential as the surrounding air. A small flame on the conductor, or some radium, may be used. The air close to the conductor is made conducting so that any charge on it leaks away and it is then at the same potential as the air near it. The potential difference between the conductor and the ground can be measured with an electrostatic voltmeter connected to the conductor and to the ground by insulated wires. If the conductor is on a pole 10 m. above the ground in the open air away from buildings or trees, the potential difference between it and the ground will be of the order of 1500 volts. The vertical field varies greatly. In fine dry weather it is usually directed downwards, indicating a negative charge on the earth's surface. It varies with the time of day and season of the year.

The vertical field has been measured at various heights by means of balloons. It is found to diminish as the height increases, and usually becomes negligible at about 10,000 m.

The air in the open is not a perfect insulator. It always contains positive and negative ions and so conducts to some extent. The conductivity can be measured by passing a stream of air through a metal case containing a charged insulated electrode, and measuring the rate at which the electrode loses its charge. Another very good method due to C. T. R. Wilson is to use a rather large horizontal metal plate placed level with the ground. The plate is insulated and connected to a capillary electrometer, which records photographically the total amount of electricity entering or leaving the plate. If the plate is covered with an earthed metal cover the charge on it is zero. If the cover is then removed, the vertical electric field induces a charge on the plate which is indicated by the electrometer, so that the strength of the field can be calculated. If the plate is left uncovered the electro-

meter indicates a small current usually flowing from the plate through the electrometer to the ground. If  $A$  is the area of the plate,  $F$  the strength of the vertical field, and  $C$  the current through the air to the plate, then  $C = \mu AF$ , where  $\mu$  is the conductivity of the air. The conductivity is of the order of  $10^{-4}$  in electrostatic units. This means that a field of 1.5 volts per centimetre gives a current of  $\frac{1}{2} \times 10^{-6}$  electrostatic units per second or  $\frac{1}{6} \times 10^{-15}$  amperes per square centimetre. If we suppose that there is a downward vertical field of 1.5 volts per centimetre all over the surface of the earth then the negative charge on the earth is about  $2.5 \times 10^{15}$  electrostatic units or one million coulombs. The total current into the earth from the air, if the conductivity were  $10^{-4}$  electrostatic units everywhere and the vertical field 1.5 volts per centimetre, would be  $2.5 \times 10^{12}$  electrostatic units or 1000 amperes, since the area of the earth's surface is  $5 \times 10^{18}$  sq. cm. The current into the earth would therefore be enough to discharge it completely in 1000 sec. or about 17 min.

## 2. How is the Earth's Charge Maintained?

One of the problems of atmospheric electricity is to explain how it is that the earth remains charged. However, the conductivity and the vertical field are not known over a sufficiently large fraction of the earth's surface for us to be sure that there is really a current of 1000 amperes going into the earth through the air. The current may be upwards over part of the surface and downwards over the rest.

There is good reason to believe that the upper regions of the atmosphere, where the pressure is low, are comparatively good conductors. It is found that electric waves are reflected from the upper regions as though there were a conducting layer at a height of about 50 Km. The earth and this conducting layer therefore form an enormous condenser, the capacity of which is about  $10^{11}$  cm. or 100,000 microfarads. If this condenser were charged so that the charge on the earth was one million coulombs the potential difference would be  $9 \times 10^6$  volts, and the vertical field would be about 2 volts per centimetre. However, the vertical field becomes small at 10 Km., so that the actual potential difference is probably usually not more than one million volts.

Various suggestions have been made to explain how the charge on the earth is maintained. C. T. R. Wilson considers that rain drops carry down enough charge to the earth to keep up its charge. Rain is usually charged more or less, and according to C. T. R. Wilson more often with negative than with positive electricity. Some observers, however, say it is generally positively charged. When a lightning flash strikes the earth it must evidently give or take away a considerable amount of electricity. We might suppose that enough flashes strike the earth on the average to keep up its charge. Very interesting observations have been made by C. T. R. Wilson, using the insulated

plate apparatus described above. When a thunderstorm occurs anywhere near this apparatus each flash is recorded by the electrometer showing a sudden rush of electricity. This indicates a sudden change in the vertical field, which then gradually changes back to its normal value. The change of field due to a discharge from a cloud to the earth can be easily calculated. Let a charge  $E$  be on a cloud at a height  $h$  above the earth. There will be an equal and opposite induced charge on the earth, and the field due to these charges is the same as that due to the charge  $E$  above the earth and a charge  $-E$  at a depth  $h$  below the surface. The two charges therefore act like a doublet of moment  $2Eh$ , and the vertical field due to this doublet on the earth's surface at a distance  $r$  is equal to  $2Eh/r^3$ , provided  $r$  is large compared with  $h$ . When the cloud is discharged by a flash to the earth the vertical field therefore suddenly changes by  $2Eh/r^3$ , so that  $Eh$  can be calculated when  $r$  is known. C. T. R. Wilson got  $r$  by measuring the time between the flash and the thunder. It was found that the results obtained could be best explained by supposing the charge  $E$  to be about 30 coulombs, and the height  $h$  from 8 to 15 Km. More flashes in which the current was upwards than downwards were observed. To keep up the negative charge on the earth, assuming it is the same all over the earth, would therefore require about 30 more upward flashes per second than downward flashes on the average. During thunderstorms the electric field is frequently directed upwards, indicating a negative charge on the clouds. A charge of 30 coulombs on a cloud is a reasonable amount of electricity. For example, a spherical cloud of radius 1 Km. with charge of 30 coulombs would have a field at its surface of only 3000 volts per centimetre, and an average density of charge of only about  $2 \times 10^{-5}$  electrostatic units per cubic centimetre. It seems to be considered that lightning flashes cannot be supposed to give more than a small fraction of the current of about 1000 amperes believed necessary to keep up the negative charge on the earth, but anyone who has seen a violent tropical thunderstorm might be inclined to doubt this opinion. For example, suppose we assume that over one-tenth of the surface of the earth there are on the average ten storms per year per hundred square miles. This gives two million storms per year. A current of 1000 amperes gives about  $3 \times 10^{10}$  coulombs in a year, so that each storm would have to give about  $2 \times 10^4$  coulombs to the earth. This would mean about 1000 flashes per storm, which seems a very low estimate. In tropical storms the flashes often follow each other so quickly that it is hard to count them, and the storm may remain in the vicinity of one place for several hours. It seems to the writer quite reasonable to suppose that lightning flashes on the average supply electricity to the earth at the rate of 1000 coulombs per second.

In thunderstorms there is usually a rapid upward current of air near the centre of the storm. The expansion of the air as it rises

causes supersaturation of the water vapour, which therefore condenses on any nuclei present. Negative ions act as nuclei more readily than positive ions, so that the drops formed will tend to be negatively charged. The drops formed fall slowly in the air, and the positive ions will therefore be carried up above the cloud into the upper conducting regions, leaving the cloud negatively charged. The positive charge which gets up to the conducting regions will be spread out to a considerable extent over the conducting layer and will induce a negative charge on the earth. If the negatively charged cloud falls far enough it may discharge to the earth either by a lightning flash or by merely carrying its charge down to the earth on the rain drops. In this way the earth gets a negative charge and the upper regions a positive charge. After the cloud is discharged to the earth, the positive charge left in the upper regions will spread out over the whole area of the conducting layer and induce an equal negative charge over the whole surface of the earth. The positive charge of course need not be carried up 50 Km. by the storm; so long as it gets up well above the clouds it will induce a negative charge in the conducting regions above it and eventually be attracted to higher levels.

Various other suggestions as to how the earth is kept charged have been made. A theory due to Ebert relies on the fact that ionized air in contact with liquid or solid bodies acquires a positive charge owing to the negative ions diffusing more rapidly than the positive. Air at the earth's surface therefore gets positively charged, and Ebert suggested that this positively charged air is carried up by air currents. This theory is not unlike the thunderstorm theory. We may also suppose that rain drops get negatively charged by the diffusion of negative ions into them and that the positive ions are carried up by air currents. Positively charged rain is frequently observed, which is a fact against the thunderstorm and Ebert's theories.

Another theory due to G. C. Simpson is that positively and negatively charged particles are shot out from the sun and that some of these reach the earth. The positive particles are supposed to be positively charged atoms which are stopped in the upper regions of the atmosphere, while the negative particles are electrons which are supposed to penetrate the atmosphere and get into the earth. This requires the electrons to have velocities almost equal to the velocity of light, and it is difficult to see how such high-velocity electrons are produced. We might suppose that they start from the sun with small velocities corresponding to the sun's temperature and are accelerated by the pressure of the sun's radiation on them, so getting nearly up to the velocity of light before they reach the earth. Such electrons might be expected to ionize the air, but W. F. G. Swann has pointed out that electrons moving with almost the velocity of light produce very few ions. They do not remain long enough near an atom to do enough

work on the electrons in the atom to knock them out. Swann has tried to detect the charges carried by these electrons by means of a large insulated mass of copper. Some of the fast electrons ought to be absorbed by such a mass so that it ought to acquire a negative charge. No such charge, however, could be detected.

Another suggestion due to W. F. G. Swann is that the very penetrating cosmic rays investigated by Millikan which come into the atmosphere from outside produce penetrating  $\beta$ -rays which travel on mainly in the direction of the cosmic rays. These  $\beta$ -rays would thus carry a negative charge along a certain distance towards the earth, and the production of such  $\beta$ -rays throughout the atmosphere all directed towards the earth would result in a flow of negative electricity into the earth from the atmosphere. If  $n$  such  $\beta$ -rays are produced per cubic centimetre per second in the atmosphere by the cosmic rays, and they all travel a distance  $l$  towards the earth, the current density due to them would be  $nle$ . The current density required to keep the earth charged is about  $\frac{1}{2} \times 10^{-6}$  electrostatic units, so that since  $e = 5 \times 10^{-10}$  we have  $\frac{1}{2} \times 10^{-6} = nl \times 5 \times 10^{-10}$  or  $nl = 10^3$ . If then  $n = 1$  we must have  $l = 10^3$  cm. These  $\beta$ -rays would ionize the air, and the number of ions per cubic centimetre would be equal to  $n$  times the number of ions produced by one  $\beta$ -ray. This, however, is a large number, since  $\beta$ -rays ionize strongly before they are stopped. It seems that this theory would give too much ionization and so will not do.

It will be seen from this discussion that we are as yet very far from having a satisfactory theory of atmospheric electricity. The facts available are not sufficient to enable definite conclusions to be reached. It is quite likely that when observations have been made all over the earth for a long time, it will appear that the observations now available were quite inadequate as a basis of a theory of the phenomena. The ionization of the atmosphere is attributed to radioactive radiations and to Millikan's cosmic rays. It is found that the air near the earth contains minute traces of radium emanation. The  $\alpha$ - and  $\beta$ -rays from this emanation and its products are sufficient to account for most of the observed ionization. The emanation is presumably evolved by traces of radium in the surface of the earth.

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## CHAPTER XVIII

# Special Relativity

### 1. Relativity in Newtonian Dynamics.

In ordinary physical experiments the building in which the experiments are made is regarded as at rest, and the positions and velocities of the bodies used are measured relatively to the building. The building is on the earth, so that we may say that the material frame of reference which is used and regarded as at rest is the earth.

All the material bodies in the building when not supported are found to have the same acceleration  $g$  vertically downwards. If, therefore, we take axes  $x, y, z$  with  $y$  vertically upwards and  $x$  and  $z$  in a horizontal plane, the equations of motion of a particle are

$$\begin{aligned}m\ddot{x} &= X, \\m\ddot{y} &= -mg + Y, \\m\ddot{z} &= Z,\end{aligned}$$

where  $x, y, z$  are the co-ordinates of the particle, and  $X, Y, Z$  are the components of any external force which may be applied to it; the axes being supposed to be fixed on the building. If  $X = Z = 0$ , and  $Y = mg$ , then  $\ddot{x} = \ddot{y} = \ddot{z} = 0$ , and the particle moves with constant velocity in a straight line.

If instead of using axes fixed on the building we use axes moving relatively to the building with constant velocity in a straight line, the equations of motion are unchanged.

For example, suppose we use axes  $x', y', z'$  fixed on an elevator in the building which is moving upwards with constant velocity  $v$ , and suppose that at time  $t = 0$  the two sets of axes coincide exactly. Then at any later time  $x = x', z = z'$ , but  $y = y' + vt$ . Substituting these values in the equations of motion, we get

$$\begin{aligned}m\ddot{x}' &= X, \\m\ddot{y}' &= -mg + Y, \\m\ddot{z}' &= Z,\end{aligned}$$

as before. If, however, we use axes moving relatively to the building with a variable velocity, the equations of the particle do not remain unchanged in form. For example, suppose we again use the axes

fixed on the elevator but that it moves upward with a constant acceleration  $a$ , then  $x = x'$ ,  $z = z'$ , and  $y = y' + \frac{1}{2}at^2$ . Substituting these values in the equations of motion, we get

$$\begin{aligned} m\ddot{x}' &= X, \\ m\ddot{y}' &= -m(g+a) + Y, \\ m\ddot{z}' &= Z. \end{aligned}$$

In this case, if  $X = 0$ ,  $Z = 0$ , and  $Y = m(g+a)$ , then  $\ddot{x}' = \ddot{y}' = \ddot{z}' = 0$ , and the particle moves with uniform velocity in a straight line. The additional force  $ma$  gives the particle an acceleration equal to that of the axes, and neutralizes the effect of the acceleration of the axes.

Let us suppose now that an observer on a material body anywhere, not necessarily on the earth, uses rectangular axes fixed on the body and observes the motion of a particle relatively to these axes. Suppose he finds the particle moves with constant velocity in a straight line when he applies no forces  $X$ ,  $Y$ ,  $Z$  to it and that in general its equations of motion are

$$m\ddot{x} = X, \quad m\ddot{y} = Y, \quad m\ddot{z} = Z.$$

What conclusions can he draw from this as to the motion of his axes through space? Is he entitled to conclude that his axes are not moving with an acceleration? As we have just seen, a motion of the axes with constant velocity in a straight line makes no difference, so it is clear that he cannot draw any conclusions as to the velocity of his system. Also, we have seen that when an acceleration is given to the axes its effect can be neutralized by applying a suitable force to the particle. It follows that the observer's axes may have any acceleration provided his system is in a field of force which neutralizes this acceleration. Thus, if his axes have an acceleration  $a$  in any direction, and if there is also present a field of force which gives to the particle an equal acceleration, then the equations of motion of the particle relative to his axes will be

$$m\ddot{x} = X, \quad m\ddot{y} = Y, \quad m\ddot{z} = Z.$$

If, for example, the observer's system is in a uniform gravitational field, then this field will give to his system and axes and to any particle he may use the same acceleration. Suppose this field is in the direction of his  $y$  axis, and that the acceleration it produces is  $f$ . Then if  $x'$ ,  $y'$ ,  $z'$  are the co-ordinates of a point in this system, and  $x$ ,  $y$ ,  $z$  the co-ordinates of the same point relative to axes at rest which coincide with  $x'$ ,  $y'$ ,  $z'$  at  $t = 0$ , we have  $x = x'$ ,  $y = y' + \frac{1}{2}ft^2$ ,  $z = z'$ , and the equations of motion relative to the fixed axes

$$m\ddot{x} = X, \quad m\ddot{y} = mf + Y, \quad m\ddot{z} = Z,$$

become

$$m\ddot{x}' = X, \quad m\ddot{y}' = Y, \quad m\ddot{z}' = Z,$$

relative to the accelerated axes.

It appears, therefore, that the observer cannot draw any conclusions as to the motion of his system through space from his observations on the motion of a particle in his system.

If the particle does not move with uniform velocity in a straight line when no forces are applied to it, this may be due either to a field of force or to a non-uniform motion of the axes, but the observer cannot tell which explanation is the correct one.

On the surface of the earth it is found that a particle, when no forces are applied to it, does not move with constant velocity but has an acceleration  $g$  vertically downwards. An observer in a laboratory may observe this acceleration  $g$ , but without other observations outside the laboratory he cannot tell whether it is due to an acceleration of the laboratory or to a field of force acting inside the laboratory or to a combination of the two. As a matter of fact, the acceleration  $g$  is believed to be partly due to the gravitational field of the earth and partly to the acceleration of the laboratory as it moves with the earth.

If there were a field of force in the laboratory which did not give the same acceleration to material particles made of different kinds of matter, such a field could be distinguished from the effect of an acceleration of the laboratory, since of course an acceleration of the laboratory must give the same opposite acceleration to any free particle in it. For example, an electric field gives no acceleration to an uncharged particle but gives one to a charged particle.

The only kind of field of force which gives the same acceleration to all particles of whatever kind is a gravitational field. We conclude, therefore, that observations on the motion of material particles can give no information as to the velocity or acceleration of the material frame of reference used, because the velocity makes no difference, and the effects of the acceleration cannot be distinguished from those of a gravitational field.

The fact that the equations of motion of a particle are unchanged by a uniform motion of the material frame of reference may be referred to as the special principle of mechanical or Newtonian relativity. According to this principle, if  $x, y, z, t$  are the co-ordinates of a particle in a system  $S$ , and  $x', y', z', t'$  the co-ordinates in another system  $S'$  moving along  $x$  with velocity  $v$  and coinciding with  $S$  at  $t = t' = 0$ , then  $x = x' + vt'$ ,  $y = y'$ ,  $z = z'$ ,  $t = t'$  are the equations for transforming the equations of motion of any particle from  $S$  to  $S'$ .

The fact that when  $S'$  has any acceleration relative to  $S$  the equations of motion in  $S'$  are still the same as in  $S$ , provided a suitable gravitational field acts in  $S'$ , may be referred to as the general principle of mechanical relativity.

## 2. The Ether, or Space.

In the study of solid geometry, the relations of the distances between different points on a solid are discussed without reference to the physical properties of the material of the solid, and in this way by a process of abstraction the idea of the geometrical properties of empty space has arisen. It has been supposed that empty space has no physical properties but only geometrical properties. No such empty space without physical properties has ever been observed, and the assumption that it can exist is without justification. It is convenient to ignore the physical properties of space when discussing its geometrical properties, but this ought not to have resulted in the belief in the possibility of the existence of empty space having only geometrical properties.

It is found that a vacuum, that is, a space from which all material particles have been removed, has important physical properties. Light waves can pass through it, and electric, magnetic, and gravitational actions can take place across it. It has specific inductive capacity and magnetic permeability. The existence of these physical properties in empty space, together with the conception of empty space having only geometrical properties, lead to the idea that space is filled with a medium, which has been called the ether, to which the physical properties in question belong. That is, we conceive really empty space as having only geometrical properties, and therefore, since we find that actual space has also physical properties, we suppose that actual space consists of our imaginary empty space filled with a medium having the observed physical properties. It is clear that there is no logical justification for this way of regarding the matter. We find that space containing no material particles has physical properties, and since we cannot separate this space in any conceivable way into two parts, one having the geometrical properties and the other the physical, we must regard the geometrical and physical properties as equally the properties of space.

If the physical properties of space could be removed, the geometrical properties would probably also disappear, in fact we are not really justified in drawing any distinction between these two sorts of properties; the geometrical properties of space are just as much physical properties as the specific inductive capacity and magnetic permeability. Descartes said that if everything inside a hollow vessel were removed its sides would be in contact. There would not even be space left in it. This profound remark shows that Descartes was not confused by an imaginary separation of the geometrical and physical properties of space.

The idea of an ether is therefore seen to be superfluous, since it is based on a purely imaginary separation of the geometrical and physical

properties of space. If it is desired to retain the word ether it may be done by using it to mean the actual space of experience as distinguished from the various imaginary types of space discussed in geometry. In what follows we shall use the word space to mean the actual space of experience with its inseparable geometrical and physical properties.

Material bodies are observed to move relatively to each other in space, and they are said to excite in the space around them fields of force which move with them. Also we have fields of force moving alone through space, without material particles moving with them. The distinction between material particles and their fields of force is not very clearly defined. The physical properties of the particles are the properties of their fields, and it is not clear that a particle is anything more than the centre from which its field radiates. The momentum of matter for example is believed to be the electromagnetic momentum of its electric and magnetic fields.

The only kinds of motion of which we have any evidence are the motions of material particles and fields of force through space. When it was supposed that space was filled with ether which was imagined to be different from space, the possibility of this ether moving through space was discussed, but we need not now consider this as a possible form of motion since we do not now admit any distinction between ether and space.

### 3. Motion through Space—the Michelson-Morley Experiment.

We may, however, consider whether the motion of a material system through space can produce any effects observable on the system. In particular, is it possible to determine the motion of the earth through space by means of observations made in a laboratory on the earth? We have seen that observations on the motion of a material particle in a laboratory give no information as to the motion of the laboratory through space. If space had only geometrical properties, we should not expect motion through it to produce any observable effects, but since this is not the case there is no *a priori* reason why motion through it should not produce effects which could be detected. We have seen that no such effects on the motion of material particles are to be expected, but the possibility of electromagnetic and optical effects remains. Up to about the year 1925 all attempts to detect any such effects failed, and it came to be generally believed that this was due, not to the experiments tried being insufficiently sensitive, but to the nature of the relations between space and matter, which were supposed to be such that the determination of the motion of a system through space by means of observations on the system was impossible. This idea was adopted by Einstein as the basis of his theory of relativity, a theory of great interest which leads to highly important results. Since 1925, however, experiments by D. C. Miller on Mount Wilson in

California, by a method originally due to Michelson, of Chicago, seemed at first sight to show that possibly optical effects due to the motion of the earth through space could be detected, and that the magnitude and direction of the velocity of the earth through space could be measured by observations made in a laboratory on the earth. These experiments threw some doubt upon the general validity of the theory of relativity, which would have had to be abandoned if Miller's results had been confirmed. The effects apparently observed by Miller were extremely small, and were most likely merely due to errors of some unexpected kind. Recent very exact repetitions of Michelson's experiment have failed to confirm Miller's results.

Let us suppose that an observer on a material system, such as the earth, which is moving through space with a velocity  $v$ , makes observations on the velocity of light. Suppose he determines the time taken by light to pass from a point  $A$  to a point  $B$  at a distance  $d$  from  $A$  and to be reflected back to  $A$ . Also, let the velocity  $v$  of his system be in the direction from  $A$  to  $B$ . Let the light start from  $A$  at time  $t_1$ , arrive at  $B$  at  $t_2$ , and get back to  $A$  at  $t_3$ . Then, if the light travels through space with velocity  $c$ , we have

$$d + v(t_2 - t_1) = c(t_2 - t_1),$$

$$d - v(t_3 - t_2) = c(t_3 - t_2),$$

or

$$t_2 - t_1 = \frac{d}{c - v},$$

$$t_3 - t_2 = \frac{d}{c + v},$$

so that

$$t_3 - t_1 = \frac{2d}{c(1 - v^2/c^2)}.$$

Suppose now that  $AB$  is at right angles to the velocity  $v$ , and that the mirror at  $B$  is arranged to reflect the light from  $A$  back to  $A$  as before. In this case  $A$  travels a distance  $v(t_3 - t_1)$  at right angles to  $AB$ , so that the distance the light goes is  $2\sqrt{d^2 + \frac{v^2}{4}(t_3 - t_1)^2}$ , and we have

$$2\sqrt{d^2 + \frac{v^2}{4}(t_3 - t_1)^2} = c(t_3 - t_1),$$

or

$$t_3 - t_1 = \frac{2d}{c\sqrt{1 - v^2/c^2}},$$

The interval  $t_3 - t_1$  is therefore not the same when  $v$  is along  $AB$  as when  $v$  is at right angles to  $AB$ , so that the observer, by measuring the time for light to go a distance  $d$  from  $A$  and back in different direc-

tions in his system, could apparently determine the magnitude and direction of the velocity of his system through space.

We have assumed here that the velocity of light  $c$  through space is a constant independent of the velocity of the material system. The source of light used moves with the system, so that we have assumed that the velocity of the light emitted by a moving source is independent of the velocity of the source. These assumptions are in accordance with the electromagnetic wave theory of light. We suppose that the velocity of light through space is equal to  $1/\sqrt{\mu K}$ , where  $\mu$  is the magnetic permeability and  $K$  the specific inductive capacity of space.

An experiment to compare the velocities of light in different directions was devised by Michelson, and carried out by him and Morley, and later by D. C. Miller.

Light from a source  $S$  (fig. 1) falls at  $45^\circ$  on a glass plate  $M$ , where it is partly reflected to a mirror  $A$  and partly transmitted to another mirror  $B$ . The mirrors  $A$  and  $B$  reflect the light back to  $M$ , where it is again partly reflected and partly transmitted. Thus two beams arrive at  $E$ , one of which has gone along the path  $SMAME$ , and the other along the path  $SMBME$ . If the two paths to  $E$  are equal, and

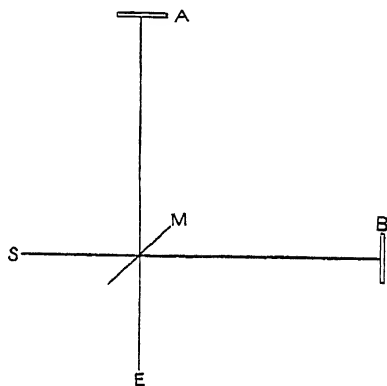


Fig. 1

white light is used, a system of interference bands is seen at  $E$ . If one of the mirrors is moved towards or away from  $M$ , the bands move. Changing  $MA$  by one-half wave-length causes a dark band to move through the distance between two adjacent dark bands. Thus, by observing the interference bands seen at  $E$ , any change in the difference between the times taken by the light to traverse the two paths can be detected. The apparatus was mounted in a horizontal plane on a block of stone floating on mercury, so that it could be slowly rotated about a vertical axis and the interference bands could be observed during the rotation. The paths  $MB$  and  $MA$  would then rotate relatively to the horizontal component of the velocity of the earth through space, and when one was parallel to this velocity component the other would be perpendicular to it. We have seen that the time for light to go a distance  $d$  and back along the direction of the motion through space with velocity  $v$  is

$$\frac{2d}{c(1 - v^2/c^2)}$$

while perpendicular to  $v$  it is

$$\frac{2d}{c\sqrt{1-v^2/c^2}},$$

so that we should expect a shift of the interference bands corresponding to a time difference

$$\frac{2d}{c} \left\{ \frac{1}{1-v^2/c^2} - \frac{1}{\sqrt{1-v^2/c^2}} \right\}$$

during the rotation of the apparatus. The bands would oscillate about a mean position twice during each revolution of the apparatus. Michelson and Morley tried this experiment with great care, and concluded that there was no appreciable effect due to the orbital motion of the earth round the sun, although they estimated that they could have detected the effect due to a velocity of one-tenth of this orbital velocity. The time difference given above is approximately equal to  $v^2d/c^3$  when  $v/c$  is small, and so corresponds to a path difference  $v^2d/c^2$  which gives a shift of the bands equal to  $v^2d/c^2\lambda$  times the distance from one band to the next one, where  $\lambda$  is the wave-length of the light used. The orbital velocity of the earth in its orbit round the sun is about  $3 \times 10^9$  cm. per second, so that if  $\lambda = 10^{-4}$  cm., we get

$$v^2d/c^2\lambda = 9 \times 10^{12} d/9 \times 10^{20} \times 10^{-4} = 10^{-4}d.$$

With  $d = 10,000$  cm., which was about the value used by Michelson and Morley,  $10^{-4}d = 1$ , so that the shift expected was about equal to the distance between two bands. No such shift was observed, and this result became the principal foundation of the theory of relativity.

#### 4. The Fitzgerald Contraction.

It was suggested by Fitzgerald and H. A. Lorentz that this negative result could be explained by supposing that material bodies contract slightly along the direction of motion when moving through space. Thus, in the Michelson-Morley experiment, if the distance along the velocity component  $v$  is  $d$ , and that perpendicular to the velocity  $d'$ , then the time difference becomes

$$\frac{2}{c} \left\{ \frac{d'}{\sqrt{1-v^2/c^2}} - \frac{d}{1-v^2/c^2} \right\},$$

which is zero if

$$d = d'\sqrt{1-v^2/c^2}.$$

Such a contraction could not be detected by ordinary measurements, provided all bodies contract equally. The Michelson-Morley experi-

ment was repeated, different materials being used to fix the distances between the mirrors, with the same negative result.

We do not know the velocity of the earth through space. We can determine the velocity of the earth relative to the sun and stars, but these bodies may be moving through space with any velocity. Suppose for example that the earth were moving through space with a velocity  $v = \frac{\sqrt{3}}{2} c$ ; then  $\sqrt{1 - v^2/c^2} = \frac{1}{2}$ , so that the length of a rod with its length perpendicular to the direction of the velocity would be double its length when parallel to the velocity.

Length therefore depends on the unknown velocity through space, and so has no absolute value, but is a relative quantity.

### 5. Einstein's Special Theory. Lorentz Transformation.

The negative result of the Michelson-Morley experiment is precisely what would have been obtained if the earth were at rest in space, and since no other effects due to the motion of the earth through space have been detected Einstein was led to put forward his *special* theory of relativity, according to which the motion of a material system through space with uniform velocity makes no difference to the phenomena observable in the system by observers on the system. This amounts to supposing that the principle of relativity, which as we have seen applies to the motion of material particles, can be extended to include electrical and optical phenomena as well.

Einstein some years later developed his *general* theory of relativity, according to which no motion of the material system could produce effects distinguishable from those due to gravitational fields either on optical phenomena or on the motion of material particles.

According to the special theory of relativity, the velocity of light should be the same in all directions to an observer on a system moving with any uniform velocity through space. Consider two such material systems  $S$  and  $S'$ , and let the velocity of  $S'$  relative to  $S$  be  $v$ . Let position and time in  $S$  be measured by co-ordinates  $x, y, z, t$  and in  $S'$  by  $x', y', z', t'$ .

Let the two sets of axes coincide at  $t = t' = 0$ , and let the relative velocity  $v$  be along the axes  $x$  and  $x'$ , so that to an observer on  $S$  the position of the origin of the co-ordinates  $x', y', z', t'$  in  $S'$  is given by  $x = vt$ , and to an observer in  $S'$  the position of the origin of  $x, y, z, t$  is given by the equation  $x' = -vt'$ .

Suppose that at the time  $t = t' = 0$ , when the two sets of axes coincide, a light wave is started at the origin of co-ordinates. To an observer on  $S$  this wave will be a sphere given by the equation  $x^2 + y^2 + z^2 = c^2 t^2$ , and to an observer on  $S'$  it will be a sphere given by  $x'^2 + y'^2 + z'^2 = c^2 t'^2$ .

There must therefore be such relations between  $x, y, z, t$  and  $x', y', z', t'$  that  $x^2 + y^2 + z^2 - c^2 t^2$  is equal to  $x'^2 + y'^2 + z'^2 - c^2 t'^2$ . To the observer on  $S$  a point at rest in  $S'$  having co-ordinates  $x, y, z$  relative to the  $S$  axes will have co-ordinates  $x - vt, y, z$  relative to the  $S'$  axes since the origin of the  $S'$  axes is at  $x = vt, y = 0, z = 0$ .

To this observer on  $S$  a measuring rod, used by an observer on  $S'$  and at rest in  $S'$ , will be moving along the  $x$  direction with velocity  $v$ , so that, assuming the Fitzgerald-Lorentz contraction, when the rod is parallel to the  $x$  axis it will be contracted, and the units of length marked on it will be lengths  $\sqrt{1 - v^2/c^2}$  instead of unity.

The observer on  $S$  will therefore consider the  $x'$  lengths as measured by the observer on  $S'$  to be measured in terms of a length  $\sqrt{1 - v^2/c^2}$  as unit, so that the  $x'$  co-ordinate of the point in question will not be  $x - vt$  when measured by the observer in  $S'$ , but  $(x - vt)/\sqrt{1 - v^2/c^2}$ . Hence

$$x' = \frac{x - vt}{\sqrt{1 - v^2/c^2}}, \quad y' = y, \quad z' = z.$$

If we substitute these values in  $x'^2 + y'^2 + z'^2 - c^2 t'^2$ , we get

$$\frac{x^2 - 2xvt + v^2 t^2}{1 - v^2/c^2} + y^2 + z^2 - c^2 t'^2,$$

which is equal to  $x^2 + y^2 + z^2 - c^2 t^2$  if

$$t' = \frac{t - \frac{v}{c^2} x}{\sqrt{1 - v^2/c^2}}.$$

Thus, according to the special principle of relativity, we have

$$y' = y, \quad z' = z, \quad x' = \frac{x - vt}{\sqrt{1 - v^2/c^2}}, \quad \text{and} \quad t' = \frac{t - \frac{vx}{c^2}}{\sqrt{1 - v^2/c^2}}.$$

In the same way, to an observer in  $S'$  a unit length along  $x$  at rest in  $S$  will be contracted to a length  $\sqrt{1 - v^2/c^2}$ , so that a point in  $S'$  at  $x', y', z'$  will have  $S$  co-ordinates given by

$$x = \frac{x' + vt'}{\sqrt{1 - v^2/c^2}}, \quad y = y', \quad z = z';$$

and this requires that

$$t = \frac{t' + \frac{vx'}{c^2}}{\sqrt{1 - v^2/c^2}}.$$

These equations giving  $x, y, z, t$  in terms of  $x', y', z', t'$  agree with those giving  $x', y', z', t'$  in terms of  $x, y, z, t$ .

## 6. Relativity of Time.

It appears that, on a material system, time as well as length is related to the velocity of the system through space, so that since the velocity through space is unknown absolute time cannot be determined. In Newtonian or classical dynamics, time was considered to be independent of position, but just how the relation between the time of an event at one place and the time of another event at another place was to be determined was not stated.

Consider a material system  $S$  moving through space with an unknown but uniform velocity, and let  $x, y, z, t$  be the space-time co-ordinates used by an observer on this system. Suppose the observer has a clock at the origin and another precisely similar clock at some other point. How can he determine the relation between the times indicated by the two clocks? Suppose a light signal sent out from the origin at a time  $t_1$  by the clock at the origin, received at the other clock at  $t_2$ , and reflected back to the origin and received there at  $t_3$ .

If the system were at rest in space we should then have  $t_2 = \frac{t_1 + t_3}{2}$ .

But according to the special principle of relativity motion through space with uniform velocity makes no difference, so Einstein considers that the relation between the times is given by  $t_2 = \frac{t_1 + t_3}{2}$  in any case, whatever the velocity of the system through space may be. That is to say, we define the relation between the times at the two clocks by the equation  $t_2 = \frac{t_1 + t_3}{2}$ . To an observer on the material

system, light travels with the same velocity in all directions, so that  $t_2$  is half-way between  $t_1$  and  $t_3$ .

In another system  $S'$  moving with uniform velocity  $v$  along the  $x$  axis of the system  $S$  and coinciding with  $S$  at  $t = t' = 0$ , an observer on  $S'$  would determine the relation between clocks in different positions in the same way as the observer in  $S$  by means of the equation  $t_2' = \frac{t_1' + t_3'}{2}$ . The times so determined in  $S'$  would not agree with

the times in  $S'$  as observed by the observer in  $S$ . To the observer in  $S$  the system  $S'$  would be moving with velocity  $v$ , so that the time  $t_2' - t_1'$  for a light signal to go from the origin of  $S'$  to a point  $(x', 0, 0)$  would not be  $x'/c$ , but would be given by  $t_2' - t_1' = \frac{x'}{c - v}$ , and the time  $t_3' - t_2'$  for the light to go back to the origin of  $S'$  would be given by

$t_3' - t_2' = \frac{x'}{c+v}$ . These equations give  $t_2' = \frac{t_3' + t_1'}{2} + \frac{v}{2c}(t_3' - t_1')$ .

Thus the assumption of the observer in  $S'$  that  $t_2' = \frac{t_3' + t_1'}{2}$  would appear to the observer on  $S$  to be in error by  $\frac{v}{2c}(t_3' - t_1')$ .

The observer on  $S'$  would consider an event happening at his origin at the time  $\frac{t_1' + t_3'}{2}$  as simultaneous with an event at  $(x', 0, 0)$  at the time  $t_2'$ , but the observer on  $S$  would consider these events not simultaneous but separated by a time interval  $\frac{v}{2c}(t_3' - t_1')$  measured in terms of the unit of time used on  $S'$ .

The assumption that  $t_2 = \frac{t_1 + t_3}{2}$  in all cases is equivalent to the assumption that light has the same velocity in all directions to an observer in any system, and, therefore, together with the Fitzgerald-Lorentz contraction, leads to the equations for  $x', y', z', t'$  in terms of  $x, y, z, t$  obtained above by making  $x^2 + y^2 + z^2 - c^2 t^2$  equal to  $x'^2 + y'^2 + z'^2 - c^2 t'^2$ .

## 7. Composition of Velocities.

The assumption that the velocity of light is the same in all directions on a system moving through space with any uniform velocity is equivalent to assuming that the vector sum of the velocity of light and any other velocity is equal to the velocity of light.

The usual rule for the composition of velocities is therefore not correct according to the special theory of relativity. Suppose we have a particle moving along the  $x$  axis with a velocity  $v$  in a system  $S$ , and another particle moving along the  $x$  axis with velocity  $u$ . We may define the difference between  $v$  and  $u$  as the velocity of the first particle to an observer moving with the second particle. Let the observer then be on a system  $S'$  moving with velocity  $u$  relatively to  $S$ . The second particle is then at rest in  $S'$ , and we may suppose that it is at the origin of  $S'$ . For the first particle let  $x = vt$ , so that for this particle in  $S'$ , since the velocity of  $S$  relative to  $S'$  is  $-u$ , we have

$$x = \frac{x' + ut'}{\sqrt{1 - u^2/c^2}}$$

and

$$t = \frac{t' + ux'/c^2}{\sqrt{1 - u^2/c^2}}.$$

Substituting these values in  $x = vt$ , we get

$$\frac{x' + ut'}{\sqrt{1 - u^2/c^2}} = \frac{vt' + \frac{vu x'}{c^2}}{\sqrt{1 - u^2/c^2}},$$

or 
$$\frac{x'}{t'} = \frac{v - u}{1 - uv/c^2}.$$

But  $x'/t'$  is the velocity of the first particle to the observer in  $S'$ , that is, by the definition adopted, the difference between  $v$  and  $u$ .

In the same way the sum of the two velocities  $u$  and  $v$  is equal to

$$\frac{v + u}{1 + uv/c^2}.$$

If  $u = c$  this becomes  $\frac{v + c}{1 + v/c} = c$ , in agreement with the original assumption that the resultant of any velocity  $v$  and the velocity of light  $c$  is equal to  $c$ .

### 8. Invariance in Expression of Physical Laws. Fizeau's Experiment.

The assumption that motion of the material system through space with any uniform velocity makes no observable difference to phenomena on the system may be expressed in another way by saying that the mathematical equations expressing physical laws in terms of the co-ordinates  $x, y, z, t$  in a system  $S$  must be identical with the equations expressing the same laws in terms of the co-ordinates  $x', y', z', t'$  in a system  $S'$  moving with any uniform velocity relative to  $S$ . The equations for  $S$  must transform into identical equations for  $S'$  when the values of  $x, y, z, t$  in terms of  $x', y', z', t'$  are substituted in them.

This principle enables the effect of the motion of matter with uniform velocity on phenomena to be determined. For example, consider the motion of light through a medium having refractive index  $\mu$ . The velocity of the light is  $c/\mu$  when the medium is at rest, so that, if the light is moving along the  $x$  axis in a system  $S$  in which the medium is at rest then we have  $x = ct/\mu$ , where  $x$  is the distance the light has travelled from the origin in the time  $t$ .

Now suppose the light is observed by an observer in another system  $S'$  moving relative to  $S$  with velocity  $-v$  along the  $x$  axis. We have

$$x = \frac{x' - vt'}{\sqrt{1 - v^2/c^2}},$$

$$t = \frac{t' - vx'/c^2}{\sqrt{1 - v^2/c^2}},$$

so that the equation  $x = ct/\mu$  becomes

$$x' - vt' = \frac{c}{\mu} (t' - vx'/c^2)$$

in the system  $S'$ . Hence

$$\frac{x'}{t'} = \frac{c/\mu + v}{1 + v/\mu c}.$$

The velocity of light in the medium moving with velocity  $v$  is therefore increased by the motion of the medium from  $c/\mu$  to  $(\frac{c}{\mu} + v)/(1 + v/\mu c)$ .

When  $v/c$  is very small this is approximately equal to

$$\frac{c}{\mu} + v \left(1 - \frac{1}{\mu^2}\right).$$

This result agrees with the experimental results of Fizeau and Michelson obtained many years before the principle of relativity was developed.

If in a system  $S$  a medium of refractive index  $\mu$  is moving along the  $x$  axis with velocity  $v$ , the velocity of light along the  $x$  axis in this medium is  $(\frac{c}{\mu} + v)/(1 + \frac{v}{\mu c})$ . The velocity of the light relative to the medium is then its velocity to an observer moving with the medium, or the difference between the two velocities, which is

$$\frac{\frac{c/\mu + v}{1 + v/\mu c} - v}{1 - \left(\frac{c/\mu + v}{1 + v/\mu c}\right) \frac{v}{c^2}} = \frac{c}{\mu},$$

according to the relativity formula for the composition of velocities.

### *Minkowski's Theory*

9. An interesting way of regarding the special principle of relativity is due to Minkowski. The phenomena which we can observe on any material system  $S$  are a succession of events, each event taking place at a definite point  $x, y, z$  and at a definite time  $t$ . An event therefore has four co-ordinates  $x, y, z, t$ . If the same event is observed on another material system  $S'$  moving relative to  $S$  with uniform velocity  $v$  along the  $x$  axis and coinciding with  $S$  at  $t = t' = 0$ , its co-ordinates in  $S'$  will be  $x', y', z', t'$  and these are related to  $x, y, z, t$  in such a way that  $x^2 + y^2 + z^2 - c^2 t^2$  is equal to  $x'^2 + y'^2 + z'^2 - c^2 t'^2$ .

Let  $\tau = ict$  and  $\tau' = ict'$ , where  $i = \sqrt{-1}$ , so that  $x^2 + y^2 + z^2 - c^2 t^2$  becomes  $x^2 + y^2 + z^2 + \tau^2$ , and  $x'^2 + y'^2 + z'^2 - c^2 t'^2$  becomes

$x'^2 + y'^2 + z'^2 + \tau'^2$ . Then if we take  $x, y, z, \tau$  to be the rectangular co-ordinates of a point in a space of four dimensions,  $\sqrt{x^2 + y^2 + z^2 + \tau^2}$  is equal to the distance of the point from the origin.

If we use another set of rectangular co-ordinates  $x', y', z', \tau'$  having the same origin but inclined to  $x, y, z, \tau$ , that is, rotate the axes  $x, y, z, \tau$  into a new position, the distance of the point  $(x, y, z, \tau)$  from the origin in terms of the new co-ordinates will be  $\sqrt{x'^2 + y'^2 + z'^2 + \tau'^2}$ , and of course this distance is unchanged by the change of axes.

In the same way, the distance between any two points  $(x_1, y_1, z_1, \tau_1)$  and  $(x_2, y_2, z_2, \tau_2)$  in the four-dimensional space is equal to

$$\sqrt{(x_2 - x_1)^2 + (y_2 - y_1)^2 + (z_2 - z_1)^2 + (\tau_2 - \tau_1)^2},$$

and this becomes

$$\sqrt{(x_2' - x_1')^2 + (y_2' - y_1')^2 + (z_2' - z_1')^2 + (\tau_2' - \tau_1')^2}$$

when the axes are changed, but its magnitude is unchanged. If we denote the distance between two points in the four-dimensional space by  $s$ , then

$$s^2 = (x_2 - x_1)^2 + (y_2 - y_1)^2 + (z_2 - z_1)^2 + (\tau_2 - \tau_1)^2,$$

and

$$ds^2 = dx^2 + dy^2 + dz^2 + d\tau^2,$$

or

$$ds^2 = dx^2 + dy^2 + dz^2 - c^2 d\tau^2,$$

where  $ds$  is the distance between two points very near together, so that  $x_2 - x_1 = dx$ , &c.  $s$  may be called the *interval* between the two events, and it is the same for all sets of rectangular axes in the four-dimensional space.

If we change from co-ordinates  $x, y, z, \tau$  to  $x', y', z', \tau'$  by rotating the axes in the plane  $(x, \tau)$  through an angle  $\theta$  from  $\tau$  towards  $x$ , then

$$x' = x \cos \theta - \tau \sin \theta,$$

$$\tau' = \tau \cos \theta + x \sin \theta,$$

$$y' = y,$$

$$z' = z.$$

Comparing with the equations

$$x' = \frac{x - vt}{\sqrt{1 - v^2/c^2}},$$

$$t' = \frac{t - vx/c^2}{\sqrt{1 - v^2/c^2}},$$

$$y' = y,$$

$$z' = z,$$

we see that  $\cos\theta = \frac{1}{\sqrt{1-v^2/c^2}}$ , and  $\tau \sin\theta = \frac{vt}{\sqrt{1-v^2/c^2}}$ , or, since  $\tau = ict$ ,

$$\sin\theta = \frac{-iv}{c\sqrt{1-v^2/c^2}}.$$

Thus we may say that changing from a system  $S$  with axes  $x, y, z, t$  to a system  $S'$  moving relative to  $S$  along  $x$  with velocity  $v$ , with axes  $x', y', z', t'$  coinciding with  $x, y, z, t$  at  $t = t' = 0$ , is equivalent to rotating the axes  $x, y, z, \tau$  through an angle  $\theta$  in the  $(x, \tau)$  plane.

The cosine of  $\theta$  is greater than unity, so that  $\theta$  is an imaginary angle.

The four-dimensional space is sometimes called the Minkowski world.

If a number of events are represented by points in this space, the configuration of these points will be fixed by the intervals  $s$  between every pair of them, and so will be the same whatever rectangular axes like  $x, y, z, \tau$  are used. The axes can be taken in any direction in the space. For example, if we draw the time axis perpendicular to the plane defined by any three points or events, these three events will be simultaneous in the resulting co-ordinates. In this way we see clearly the relative character of space and time, but it appears that there are absolute quantities connected with events, namely the intervals  $s$  between any two of them.

A curve drawn in the four-dimensional space is called a world line, and the world line of a particle represents the relation between its position and the time in any set of co-ordinates which may be selected. The shape of the world line is the same for any set of co-ordinates.

Since the configuration of the points representing events in the four-dimensional space represents what happens in the system considered, and is independent of the co-ordinate system used, the laws according to which the events happen should be capable of being expressed in a form independent of the co-ordinate system chosen. Any vector quantity should be capable of representation by a line drawn in the four-dimensional space, and so should have four components in any particular system of co-ordinates  $x, y, z, \tau$ . The length of the line representing the vector quantity will be the same for all co-ordinate systems, and in any co-ordinate system its direction will be related in the same way to the world lines and points which represent the events in the four-dimensional space. The interval  $s$  between two events is an example of such a four-dimensional vector.

The components of such a vector will be obtained by multiplying its length by the cosines of the angles it makes with the axes. Let  $a_1, a_2, a_3, a_4$  be these cosines for an element  $ds$  of a world line. The

components of  $ds$  are  $dx$ ,  $dy$ ,  $dz$ , and  $d\tau$ , so that  $\alpha_1 = dx/ds$ ,  $\alpha_2 = dy/ds$ ,  $\alpha_3 = dz/ds$ , and  $\alpha_4 = d\tau/ds$ . But  $ds^2 = dx^2 + dy^2 + dz^2 - c^2 d\tau^2$ , and

$$v^2 = \left(\frac{dx}{dt}\right)^2 + \left(\frac{dy}{dt}\right)^2 + \left(\frac{dz}{dt}\right)^2,$$

so that  $\left(\frac{ds}{dt}\right)^2 = v^2 - c^2$ , or  $\frac{dt}{ds} = \frac{-i}{\sqrt{c^2 - v^2}}$ , where  $i = \sqrt{-1}$ .

Therefore

$$\alpha_1 = \frac{dx}{ds} = \frac{dx}{dt} \frac{dt}{ds} = \frac{-iv_x}{\sqrt{c^2 - v^2}},$$

and in the same way

$$\alpha_2 = \frac{-iv_y}{\sqrt{c^2 - v^2}}, \quad \alpha_3 = \frac{-iv_z}{\sqrt{c^2 - v^2}},$$

and

$$\alpha_4 = \frac{d\tau}{ds} = ic \frac{dt}{ds} = \frac{c}{\sqrt{c^2 - v^2}}.$$

In this way we can express the direction cosines of  $ds$  in terms of the velocity  $v$  corresponding to any particular set of co-ordinates  $x, y, z, \tau$ .

## 10. Minkowski Velocity.

Let us now consider velocity in the Minkowski world. We wish to find a four-dimensional vector which is independent of the particular co-ordinates chosen, and which may be taken to represent this velocity. Consider an element  $ds$  of the world line of a particle, and let the  $\tau$  axis be drawn along  $ds$ . In this case  $ds = d\tau = icdt$ , so that  $ds/dt = ic$  is the velocity of the particle along its world line. Let us suppose then that the Minkowski velocity of a particle is equal to  $ic$  in any co-ordinates, and directed along  $ds$ . When the  $\tau$  axis does not coincide with  $ds$  then the velocity  $ic$  has components  $ic\alpha_1, ic\alpha_2, ic\alpha_3$ , and  $ic\alpha_4$ , which are equal to

$$\frac{v_x}{\sqrt{1 - v^2/c^2}}, \quad \frac{v_y}{\sqrt{1 - v^2/c^2}}, \quad \frac{v_z}{\sqrt{1 - v^2/c^2}}, \quad \text{and} \quad \frac{ic}{\sqrt{1 - v^2/c^2}},$$

When  $v/c$  is very small, these are approximately

$$v_x, v_y, v_z, \text{ and } ic.$$

Thus the  $x, y, z$  components of the Minkowski velocity  $ic$  are equal to the corresponding components of the three-dimensional velocity  $v$  when  $v/c$  is very small.

It is important to note that the Minkowski velocity vector  $ic$  directed along  $ds$  is only a velocity in the sense that, when a set of axes  $x, y, z, \tau$  is chosen, the components of  $ic$  are velocities relative to the axes chosen. It is only when axes have been chosen that the position of the particle can be specified. If we choose a set of axes, the co-ordinates  $(x, y, z, \tau)$  of any point on the world line give the position of the particle  $(x, y, z)$  at the time  $t$  given by  $\tau = ict$ , and its velocity is equal to  $ic$  directed along  $ds$  relative to the axes chosen. The velocity is the same for any set of axes, but in the absence of axes  $(x, y, z, \tau)$  there is no velocity.

### 11. Minkowski Force.

If the world line of a particle is straight, its velocity is constant in magnitude and direction, so that there is no force acting on it in any set of axes. If the world line is curved, the direction of the Minkowski velocity  $ic$  along  $ds$  changes with the time, so that a force must be acting on the particle. Let the direction of  $s$  change through an angle  $d\theta$  in a time  $dt$ , and let  $ds$  be the element of  $s$  described in the time  $dt$ . Since the Minkowski velocity  $ic$  is constant along  $ds$ , its change in  $dt$  is equal to  $icd\theta$ , and is perpendicular to  $ds$ . The acceleration is therefore equal to  $icd\theta/dt$ , which is equal to  $ic \frac{d\theta}{ds} \frac{ds}{dt}$ . If  $r$  is the radius of curvature of the

world line, we have  $rd\theta = ds$ , and also  $\frac{ds}{dt} = \frac{ic}{\beta}$ , where  $\beta = \frac{1}{\sqrt{1-v^2/c^2}}$ , so that

$$ic \frac{d\theta}{dt} = \frac{ic}{r} \frac{ic}{\beta} = -\frac{c^2}{r\beta}.$$

In classical dynamics the force on a particle is equal to its mass multiplied by its acceleration. We suppose that this is true when the velocity of the particle is very small compared with the velocity of light, so that  $\beta = \frac{1}{\sqrt{1-v^2/c^2}} = 1$ . In

this case the acceleration  $ic \frac{d\theta}{dt} = -\frac{c^2}{r}$ , and we may therefore define the Minkowski force, which will be denoted by  $P$ , by the equation  $P = -mc^2/r$ . It is directed along the radius of curvature  $r$  towards the centre of curvature. When  $\beta$  is not equal to unity, the acceleration  $icd\theta/dt$ , which will be denoted by  $a$ , is given by  $a = -c^2/r\beta$ , so that

$$P = \frac{-mc^2}{r} = m\beta a.$$

The components of the Minkowski velocity are  $\beta v_x$ ,  $\beta v_y$ ,  $\beta v_z$ , and  $ic\beta$ , so that the components of  $a$  are  $\frac{d}{dt}(\beta v_x)$ ,  $\frac{d}{dt}(\beta v_y)$ ,  $\frac{d}{dt}(\beta v_z)$ , and  $\frac{d}{dt}(ic\beta)$ .

The components of the Minkowski force  $P$  are therefore

$$\begin{aligned} P_x &= m\beta \frac{d}{dt}(\beta v_x), & P_y &= m\beta \frac{d}{dt}(\beta v_y), \\ P_z &= m\beta \frac{d}{dt}(\beta v_z), & P_\tau &= m\beta \frac{d}{dt}(ic\beta) = icm\beta \frac{d\beta}{dt}. \end{aligned}$$

Now  $\frac{d\beta}{dt} = \frac{v}{c^2} \beta^3 \frac{dv}{dt}$ , where  $\frac{dv}{dt}$  is simply the rate of increase of the magnitude of  $v$  without reference to direction, and so not equal to the acceleration. When  $v=0$  and therefore  $\beta=1$ , this makes  $\frac{d\beta}{dt}=0$ , and the components of  $P$  are then

$$\begin{aligned} P_x &= m \frac{dv_x}{dt}, & P_y &= m \frac{dv_y}{dt}, \\ P_z &= m \frac{dv_z}{dt}, & P_\tau &= 0, \end{aligned}$$

as in classical dynamics.

We now define a force  $F$  by the equation  $F = ma$ , which agrees with the classical definition when  $v = 0$  and  $\beta = 1$ ; then since  $P = m\beta a$  we have  $P = \beta F$ . The components of  $F$  are then

$$\begin{aligned} F_x &= m \frac{d}{dt} (\beta v_x), & F_y &= m \frac{d}{dt} (\beta v_y), \\ F_z &= m \frac{d}{dt} (\beta v_z), & F_\tau &= icm \frac{d\beta}{dt}. \end{aligned}$$

If we put  $m\beta = m'$  these equations become

$$\begin{aligned} F_x &= \frac{d}{dt} (m' v_x), & F_y &= \frac{d}{dt} (m' v_y), \\ F_z &= \frac{d}{dt} (m' v_z), & F_\tau &= ic \frac{dm'}{dt}, \end{aligned}$$

since  $m$  is a constant. Thus it appears that the mass of a particle, defined as usual by the equation  $F_x = \frac{d}{dt} (m' v_x)$ , is not a constant, but is equal to  $\frac{m}{\sqrt{1-v^2/c^2}}$ , where  $m$  is its mass when  $v/c$  is very small.

This variation of mass with velocity has been verified for electrons by experiments on the  $\beta$ -rays of radium and on cathode rays (see Chap. IX, sections 7 and 8).

The force  $F$  is not a four-dimensional vector, since its magnitude is equal to  $-\frac{mc^2}{r\beta}$  and so depends on  $v$  and therefore on the particular set of axes  $x, y, z, \tau$  chosen. The Minkowski force  $P = -mc^2/r$  in any set of axes and so is a four-dimensional vector.

Two particular cases will now be considered. In one the force  $F$  is perpendicular to the velocity  $v$ , and in the other  $F$  is along the direction of  $v$ .  $P$  and  $F$  are at right angles to  $ds$ , and  $v$  lies in the plane containing  $ds$  and the  $\tau$  axis. Hence, when  $F$  is perpendicular to  $v$  it is perpendicular to the  $\tau$  axis, so that  $F_\tau = 0$  and consequently  $\frac{d\beta}{dt} = 0$ ,  $v$  being thus a constant, as we should expect.  $a_\tau = \frac{d}{dt} (ic\beta)$  is also zero. In this case, therefore, since  $\beta$  is constant,

$$\begin{aligned} F_x &= m' \frac{dv_x}{dt}, & F_y &= m' \frac{dv_y}{dt}, \\ F_z &= m' \frac{dv_z}{dt}, & F_\tau &= 0, \end{aligned}$$

so that  $F = m'f$ , where  $f$  is the three-dimensional acceleration relative to  $x, y, z$ . The particle therefore behaves as if it had a constant mass  $m' = \frac{m}{\sqrt{1-v^2/c^2}}$ .

When  $F$  is along  $v$ , it is at right angles to  $\tau$ , and in the plane containing  $ds$  and  $\tau$ . But  $F$  and  $P$  are always perpendicular to  $ds$ , so that in this case  $ds$  must coincide with the  $\tau$  axis and  $v$  must be 0. In this case, therefore,

$$F_x = m \frac{d}{dt} (\beta v_x) = m v_x \frac{d\beta}{dt} + m\beta \frac{dv_x}{dt},$$

which, when  $v = 0$ , reduces to  $F_x = m \frac{dv_x}{dt}$ .

In the same way  $F_y = m \frac{dv_y}{dt}$ ,  $F_z = m \frac{dv_z}{dt}$ , and  $F_\tau = 0$ .

Thus when  $F$  is along  $v$  the mass of the particle is equal to  $m$ , its mass when at rest, because in this case  $v$  must be zero.

Another particular case is that in which  $F$  lies in the plane containing  $ds$ ,  $\tau$ , and  $v$ . In this case, without loss of generality we may take the  $x$  axis along  $v$ , so that the equations for the components of  $F$  become

$$F_x = F_v = m \frac{d}{dt}(\beta v), \quad F_y = 0, \quad F_z = 0, \quad F_\tau = icm \frac{d\beta}{dt},$$

since here  $v_x = v$ .

The force  $F$  is at right angles to  $ds$ , and  $F_v$  is at right angles to  $\tau$ , so that the angle between  $F$  and  $F_v$  is equal to the angle between  $ds$  and  $\tau$ , the cosine of which is  $\beta$ .

Hence  $F_v = F\beta$ . Also  $\frac{d}{dt}(\beta v) = \beta^3 \frac{dv}{dt}$ , so that  $F_v = m\beta^3 \frac{dv}{dt}$ , and  $F = m\beta^2 \frac{dv}{dt}$ .

The particle moves along a straight line with velocity  $v$  and acceleration  $dv/dt$ , so that since  $F_v$  is the component of  $F$  along the line it has been said to behave as if it had a mass equal to  $m\beta^3$ . However, since  $F_v = \frac{d}{dt}(m\beta v) = \frac{d}{dt}(m'v)$  it is better to regard the particle as behaving as if it had a mass equal to  $m' = \beta m$ .

## 12. Work and Energy in Minkowski World.

Let us now consider work and energy in the four-dimensional world. Since  $P$  and  $F$  are both at right angles to  $ds$  we have

$$F_x dx + F_y dy + F_z dz + F_\tau d\tau = 0,$$

where  $dx$ ,  $dy$ ,  $dz$ , and  $d\tau$  are the components of  $ds$ . If we denote the work by  $W$ , and define it by the equation

$$dW = F_x dx + F_y dy + F_z dz,$$

we get

$$dW + F_\tau d\tau = 0.$$

In the case of a force  $F$  acting on a particle of mass  $m$ , we have  $F_\tau = icm \frac{d\beta}{dt}$  and  $d\tau = icdt$ , so that  $dW = mc^2 d\beta$ . Hence  $W = mc^2 \beta + C$ .

To determine the constant  $C$ , let  $W = 0$  when  $\beta = 1$ , so that  $C = -mc^2$ . Hence  $W = mc^2(\beta - 1)$ . When  $\beta = 1$ ,  $v = 0$ , so that  $W$  is the kinetic energy of the particle. When  $v/c$  is very small,  $\beta = \frac{1}{\sqrt{1 - v^2/c^2}} = 1 + \frac{v^2}{2c^2}$ , so that we get  $W = mv^2/2$ , as in classical dynamics.

Now

$$m' = m\beta \quad \text{so that} \quad W = c^2(m' - m).$$

This shows that when the energy of the particle is increased by  $W$ , its mass is increased by  $W/c^2$ . Thus it appears that kinetic energy has mass equal to the energy divided by the square of the velocity of light.

We have so far regarded the mass  $m$  of the particle considered when at rest as a constant, and we defined the Minkowski force by the equation  $P = -mc^2/r$ . When  $m$  is supposed to vary, and changes to  $m + dm$  when  $ds$  turns through an angle  $d\theta$ , the momentum  $mic$  along  $ds$  changes by  $mic d\theta$  at right angles to  $ds$ , and  $icdm$  along  $ds$ . Thus, in axes relatively to which the particle is at rest, we get a component of  $P$  along  $ds$  equal to  $ic \frac{dm}{dt}$ , which is equal to  $-c^2 \frac{dm}{ds}$ , since  $ds = icdt$  when  $\beta = 1$ .

The Minkowski force therefore consists of a component  $-mc^2/r$  at right angles to  $ds$ , and a component  $P_s = -c^2 \frac{dm}{ds}$  along  $ds$ . Now  $-c^2 \frac{dm}{ds} = -c^2 \frac{dm}{dt} \frac{dt}{ds}$  so that, since  $\frac{dt}{ds} = -\frac{i}{c}\beta$ , we get  $-c^2 \frac{dm}{ds} = ic\beta \frac{dm}{dt}$  in any set of axes.

The Minkowski work may be defined as the negative product of the Minkowski force along  $ds$  into  $ds$  or  $-P_s ds$ . The component of  $P$  perpendicular to  $ds$  then does no work. We have  $-P_s ds = +c^2 dm$ . Now we have seen that when  $m$  is constant then  $c^2(m' - m)$  is equal to the kinetic energy, which suggests that energy has mass equal to the energy divided by  $c^2$ . This result agrees with the equation  $-P_s ds = c^2 dm$ , if we admit that the Minkowski work represents energy  $E$  supplied to the particle. We may then put  $dE = -P_s ds = c^2 dm$ , so that, if  $m = 0$  when  $E = 0$ , we have  $E = mc^2$ . The Minkowski force was defined as rate of change of momentum, and the Minkowski work must therefore be of the nature of energy. The negative sign which has to be given to the product is not surprising in view of the fact that  $P_s = -c^2 \frac{dm}{ds}$ .

The theory of relativity therefore leads to the very important conclusion that a body of mass  $m$  has energy  $mc^2$  in virtue of its mass  $m$ . The energy given by  $E = mc^2$  may be called the Minkowski energy, since it is independent of the axes chosen.

The Minkowski work is not the same as ordinary work unless  $\beta = 1$  and  $\frac{dm}{dt} = 0$ ; for ordinary work  $W$  was defined by  $dW = F_x dx + F_y dy + F_z dz$  and  $P_x = \beta F_x$ ,  $P_y = \beta F_y$ ,  $P_z = \beta F_z$ ,  $P_t = \beta F_t$ , and consequently Minkowski work is equal to ordinary work when  $m$  remains constant and  $\beta = 1$ , because then  $P = 0$ . The Minkowski energy  $mc^2$  is therefore also equal to ordinary energy when  $\frac{dm}{dt} = 0$  and

$v = 0$ . A particle at rest therefore has ordinary energy  $mc^2$ . When moving, its kinetic energy is  $c^2(m' - m)$ , and its total ordinary energy  $c^2m'$ , but its Minkowski energy is still  $c^2m$ . Hence if  $E$  denotes the Minkowski energy and  $E'$  the ordinary energy then  $E' = \beta E$ .

The components of  $P_s = ic\beta \frac{dm}{dt}$  along the axes  $x, y, z, \tau$  are  $\beta^2 v_x \frac{dm}{dt}, \beta^2 v_y \frac{dm}{dt}, \beta^2 v_z \frac{dm}{dt}$ , and  $ic\beta^2 \frac{dm}{dt}$ .

Hence 
$$P_x = m\beta \frac{d}{dt}(\beta v_x) + \beta^2 v_x \frac{dm}{dt},$$

or 
$$P_x = \beta \frac{d}{dt}(m\beta v_x),$$

with similar equations for  $P_y$  and  $P_z$ , and  $P_\tau = ic\beta \frac{d}{dt}(m\beta)$ . Since  $\tau = ict$ , we may define the  $t$  component of  $P$  by the equation  $P_\tau = iP_t$ , so that  $P_t = c\beta \frac{d}{dt}(m\beta)$ . The components of the force  $F$  defined by  $P = \beta F$  are therefore

$$F_x = \frac{d}{dt}(m'v_x), \quad F_y = \frac{d}{dt}(m'v_y),$$

$$F_z = \frac{d}{dt}(m'v_z), \quad F_t = c \frac{dm'}{dt},$$

where  $m' = m\beta$ , as before, but  $m$  is not now supposed to be constant.

For a particle moving with a constant velocity  $v$ , we have  $F_x = v_x \frac{dm'}{dt}$ , with similar equations for  $F_y$  and  $F_z$ , so that if  $F_\tau$  denotes the component of  $F$  along the direction of  $v$ , which is perpendicular to the  $\tau$  axis,  $F_v = v \frac{dm'}{dt}$ . Thus there is a three-dimensional force  $F$  on the particle just sufficient to give it the momentum required to keep its velocity constant.

As an example consider a star like the sun which is radiating energy into the surrounding space. In the case of the sun the mass of the energy radiated is about four million tons per second. If the star is moving with uniform velocity, there must be a force on it equal to  $v dm'/dt$ . This force is produced by the flow of energy away from the star, which merely means that the energy radiated has momentum equal to that lost by the star, so that the velocity of the star remains constant because the momentum lost is proportional to the mass lost.

It appears that according to the theory of relativity the mass of any system is equal to its energy divided by  $c^2$ . The principles of the conservation of mass and of energy are therefore identical. The

energy changes which occur in laboratory experiments are too small to produce appreciable changes of mass. For example, 1 gm. of radium evolves about 100 calories of heat energy per hour. The mass of this energy evolved in one year is therefore  $(365 \times 24 \times 100 \times 42 \times 10^6) \div (9 \times 10^{20})$  gm., which is only about 1/20,000 of a milligram.

The helium atom is believed to be made up of four hydrogen atoms, but its atomic weight is 4 whereas that of hydrogen is 1.007. It is therefore supposed that when hydrogen atoms combine to form 4 gm. of helium, an amount of energy equal to  $0.028 \times 9 \times 10^{20} = 2.5 \times 10^{19}$  ergs must be evolved.

### 13. World Tensors.

Consider any two four-dimensional vectors  $p$  and  $q$ , having components  $p_x, p_y, p_z, p_\tau = ip_t$ , and  $q_x, q_y, q_z, q_\tau = iq_t$ . We can form sixteen products of one component of  $p$  with one of  $q$  thus:

$$\begin{array}{cccc} p_x q_x & p_x q_y & p_x q_z & ip_x q_t \\ p_y q_x & p_y q_y & p_y q_z & ip_y q_t \\ p_z q_x & p_z q_y & p_z q_z & ip_z q_t \\ ip_t q_x & ip_t q_y & ip_t q_z & -p_t q_t. \end{array}$$

These sixteen products are said to be the sixteen components of a world tensor. In general a world tensor is defined as a quantity having sixteen components which transform from one set of axes  $x, y, z, \tau$  to any other set in the same way as the sixteen products of the components of a pair of vectors.

If  $t$  denotes a world tensor, its components may be denoted by  $t$  with two suffixes, as, for example,  $t_{xy}$  or  $t_{y\tau}$ .

If interchanging the two suffixes does not change the value of any component so that, for example,  $t_{xy} = t_{yx}$  and  $t_{y\tau} = t_{\tau y}$ , the tensor is said to be symmetrical, and can be specified by ten quantities.

If interchanging the two suffixes of any component merely changes the sign of the component so that, for example,  $t_{xy} = -t_{yx}$  and  $t_{z\tau} = -t_{\tau z}$ , then  $t_{xx}, t_{yy}, t_{zz}$ , and  $t_{\tau\tau}$  are all zero, and the tensor is said to be skew-symmetrical, and can be specified by only six quantities, viz.  $t_{xy}, t_{xz}, t_{yz}, t_{x\tau}, t_{y\tau}, t_{z\tau}$ .

The vector divergence  $\mathbf{V}$  of a world tensor is defined by the equations giving its four components. The  $x$  component is given by

$$V_x = \frac{\partial t_{xx}}{\partial x} + \frac{\partial t_{xy}}{\partial y} + \frac{\partial t_{xz}}{\partial z} + \frac{\partial t_{xt}}{c \partial t},$$

where  $\frac{\partial t_{xt}}{c \partial t} = \frac{\partial t_{x\tau}}{\partial \tau}$ , since  $d\tau = icdt$  and  $t_{x\tau} = it_{xt}$ , with similar equations for  $V_y, V_z$ , and  $V_\tau = iV_t$ .

The components of the vector divergence of a skew-symmetrical tensor are then given by the equations

$$\begin{aligned} V_x &= \frac{\partial t_{xy}}{\partial y} + \frac{\partial t_{xz}}{\partial z} + \frac{\partial t_{xt}}{\partial t}, \\ V_y &= \frac{\partial t_{yx}}{\partial x} + \frac{\partial t_{yz}}{\partial z} + \frac{\partial t_{yt}}{\partial t}, \\ V_z &= \frac{\partial t_{zx}}{\partial x} + \frac{\partial t_{zy}}{\partial y} + \frac{\partial t_{zt}}{\partial t}, \\ V_t &= \frac{\partial t_{tx}}{\partial x} + \frac{\partial t_{ty}}{\partial y} + \frac{\partial t_{tz}}{\partial z}. \end{aligned}$$

Now let  $H_x = t_{yz}$ ,  $H_y = t_{zx}$ , and  $H_z = t_{xy}$ ; and also let  $E_x = t_{tx}$ ,  $E_y = t_{ty}$ ,  $E_z = t_{tz}$ ; so that

$$\begin{aligned} V_x &= \frac{\partial H_z}{\partial y} - \frac{\partial H_y}{\partial z} - \frac{1}{c} \frac{\partial E_x}{\partial t}, \\ V_y &= \frac{\partial H_x}{\partial z} - \frac{\partial H_z}{\partial x} - \frac{1}{c} \frac{\partial E_y}{\partial t}, \\ V_z &= \frac{\partial H_y}{\partial x} - \frac{\partial H_x}{\partial y} - \frac{1}{c} \frac{\partial E_z}{\partial t}, \\ V_t &= \frac{\partial E_x}{\partial x} + \frac{\partial E_y}{\partial y} + \frac{\partial E_z}{\partial z}, \end{aligned}$$

since  $t_{yz} = -t_{zy}$ ,  $t_{tz} = -t_{zt}$ , &c.

Thus if  $\mathbf{V}_1$  denotes the component of  $\mathbf{V}$  at right angles to the  $\tau$  axis, so that  $V^2 = V_1^2 + V_\tau^2$  and  $V_1^2 = V_x^2 + V_y^2 + V_z^2$ , and if  $\mathbf{E}$ ,  $\mathbf{H}$  denote the three-dimensional vectors of which the components are  $E_x$ ,  $E_y$ ,  $E_z$ , and  $H_x$ ,  $H_y$ ,  $H_z$ , we have

$$\mathbf{V}_1 + \frac{1}{c} \frac{\partial \mathbf{E}}{\partial t} = \text{curl } \mathbf{H}$$

and

$$V_t = \text{div } \mathbf{E}.$$

In the four-dimensional space the curl or rotation of a vector may be defined just as in three-dimensional space. Thus the components of the curl  $\mathbf{R}$  of a vector  $\mathbf{P}$  having components  $P_x$ ,  $P_y$ ,  $P_z$ ,  $P_\tau = iP_t$  are defined by the equations

$$\begin{aligned} R_{xy} &= \frac{\partial P_y}{\partial x} - \frac{\partial P_x}{\partial y}, & R_{yz} &= \frac{\partial P_z}{\partial y} - \frac{\partial P_y}{\partial z}, \\ R_{zx} &= \frac{\partial P_x}{\partial z} - \frac{\partial P_z}{\partial x}, & R_{x\tau} &= \frac{\partial P_\tau}{\partial x} - \frac{\partial P_x}{\partial \tau}, \\ R_{y\tau} &= \frac{\partial P_\tau}{\partial y} - \frac{\partial P_y}{\partial \tau}, & R_{z\tau} &= \frac{\partial P_\tau}{\partial z} - \frac{\partial P_z}{\partial \tau}. \end{aligned}$$

Differentiating  $R_{xy}$  with respect to  $z$ ,  $R_{yz}$  with respect to  $x$ , and  $R_{zx}$  with respect to  $y$ , and adding the three equations so obtained, we get

$$\frac{\partial R_{yz}}{\partial x} + \frac{\partial R_{zx}}{\partial y} + \frac{\partial R_{xy}}{\partial z} = 0.$$

In the same way we get

$$\frac{\partial R_{z\tau}}{\partial y} + \frac{\partial R_{\tau y}}{\partial z} + \frac{\partial R_{yz}}{\partial \tau} = 0,$$

$$\frac{\partial R_{\tau x}}{\partial z} + \frac{\partial R_{xz}}{\partial \tau} + \frac{\partial R_{z\tau}}{\partial x} = 0,$$

$$\frac{\partial R_{xy}}{\partial \tau} + \frac{\partial R_{y\tau}}{\partial x} + \frac{\partial R_{\tau x}}{\partial y} = 0.$$

We have  $R_{xy} = -R_{yx}$ , &c., so that the six components of the curl  $\mathbf{R}$  may be regarded as the components of a skew-symmetrical world tensor. Thus when a world tensor is the curl of a vector it must be a skew-symmetrical tensor, and its components must satisfy the four equations just obtained.

Let us now suppose that the world tensor  $\mathbf{R}$  is identical with the tensor  $\mathbf{t}$  previously considered, so that

$$R_{yz} = H_x, \quad R_{zx} = H_y, \quad R_{xy} = H_z,$$

$$R_{tx} = E_x, \quad R_{ty} = E_y, \quad R_{tz} = E_z,$$

where

$$R_{\tau x} = iR_{tx}, \quad R_{\tau y} = iR_{ty}, \quad R_{\tau z} = iR_{tz}.$$

Substituting these values we obtain

$$\frac{\partial H_x}{\partial x} + \frac{\partial H_y}{\partial y} + \frac{\partial H_z}{\partial z} = 0,$$

$$-\frac{\partial E_z}{\partial y} + \frac{\partial E_y}{\partial z} - \frac{1}{c} \frac{\partial H_x}{\partial t} = 0,$$

$$\frac{\partial E_x}{\partial z} + \frac{1}{c} \frac{\partial H_y}{\partial t} - \frac{\partial E_z}{\partial x} = 0,$$

$$-\frac{1}{c} \frac{\partial H_z}{\partial t} - \frac{\partial E_y}{\partial x} + \frac{\partial E_x}{\partial y} = 0,$$

or

$$\operatorname{div} \mathbf{H} = 0, \quad \text{and} \quad \operatorname{curl} \mathbf{E} = -\frac{1}{c} \frac{\partial \mathbf{H}}{\partial t}.$$

Thus it appears that if we have any skew-symmetrical world tensor  $\mathbf{t}$  which is equal to the curl of a four-dimensional vector  $\mathbf{P}$ , and if we put

$$t_{yz} = H_x, \quad t_{zx} = H_y, \quad t_{xy} = H_z,$$

$$t_{tx} = E_x, \quad t_{ty} = E_y, \quad t_{tz} = E_z,$$

we shall have

$$\begin{aligned} \operatorname{div} \mathbf{H} &= 0, \quad \operatorname{div} \mathbf{E} = V_t, \\ \operatorname{curl} \mathbf{H} &= \mathbf{V}_1 + \frac{1}{c} \frac{\partial \mathbf{E}}{\partial t}, \quad \operatorname{curl} \mathbf{E} = -\frac{1}{c} \frac{\partial \mathbf{H}}{\partial t}, \end{aligned}$$

where  $V_t$  is the time component of the vector divergence of the tensor  $t$  and  $\mathbf{V}_1$  is the component perpendicular to the time axis. These results of course are true for any set of axes  $x, y, z, \tau$  in the four-dimensional world.

#### 14. The Electromagnetic Equations.

Maxwell's equations of the electromagnetic field in the form given to them by H. A. Lorentz are (see Chapter I):

$$\begin{aligned} \operatorname{div} \mathbf{H} &= 0, \quad \operatorname{div} \mathbf{E} = \rho, \\ \operatorname{curl} \mathbf{H} &= \frac{1}{c} \left( \rho \mathbf{v} + \frac{\partial \mathbf{E}}{\partial t} \right), \quad \operatorname{curl} \mathbf{E} = -\frac{1}{c} \frac{\partial \mathbf{H}}{\partial t}, \end{aligned}$$

where  $\mathbf{H}$  is the magnetic field strength,  $\mathbf{E}$  the electric field strength,  $\rho$  the density of electricity or charge per unit volume, and  $\mathbf{v}$  the velocity of the electricity. Comparing these equations with those just obtained above, we see that if  $V_t = \rho$  and  $\mathbf{V}_1 = \rho \mathbf{v}/c$  the two sets of equations become identical.

If  $e$  is the charge in a volume  $S$  which is at rest relatively to the axes used, so that  $\rho = e/S$ , and if we change to axes in which the charge is moving with velocity  $v$ , the volume changes to  $S/\beta$  so that the density of charge becomes  $e\beta/S$ . Hence, if we put  $\rho' = \rho/\beta$ ,  $\rho'$  has the same value for any set of axes.  $\rho'$  may be called the Minkowski density of charge.

The Minkowski current density may be defined as the product of  $\rho'$  and the Minkowski velocity, so that it is equal to  $\rho'ic$ , and its components are  $\rho'\beta v_x = \rho v_x$ ,  $\rho v_y$ ,  $\rho v_z$ , and  $ic\rho$ . Its time component is therefore  $\rho c$ , and its component perpendicular to the  $t$  axis is  $\rho \mathbf{v}$ . If then  $V_t = \rho$  and  $\mathbf{V}_1 = \rho \mathbf{v}/c$ , it appears that  $\mathbf{V}$  is equal to the Minkowski current divided by  $c$ .

Maxwell's electromagnetic equations therefore show that the three components of the magnetic field and the three components of the electric field may be regarded as the six components of a skew-symmetrical world tensor, which is called the electromagnetic field tensor. This tensor is equal to the curl of a world vector, which may be called the Minkowski vector potential, and its vector divergence is equal to the Minkowski current divided by the velocity of light.

As an example, suppose that we have a charge  $e$  in a magnetic field  $H = H_y$  along the  $y$  axis, and suppose that the charge is at rest relative to the axes. If now we change to axes in which the charge has a velocity  $v_x$  along the  $x$  axis, we are rotating the axes in the  $(x, \tau)$

plane through an angle the cosine of which is  $\beta$ . Then  $H_y = t_{zx}$ , and so  $H_y$  transforms in the same way as the product of two vectors, one along the  $z$  axis and one along the  $x$  axis. The rotation in the  $(x, \tau)$  plane leaves a vector along  $z$  unchanged in position, but a vector along  $x$  is changed into one inclined to the  $x$  axis in the  $(x, \tau)$  plane, and its components are proportional to  $\beta$  along  $x$  and  $iv_x\beta/c$  along  $\tau$ . Thus the magnetic field  $H_y$  transforms into a magnetic field  $\beta H_y$  along  $y$  and an electric field  $E_z = v_x\beta H_y/c$  along  $z$ , because  $E_z = t_{tz}$ . Thus when the charge  $e$  is moving with velocity  $v_x$  along the  $x$  axis in a magnetic field of strength  $\beta H_y$  along the  $y$  axis, there is a force on it along the  $z$  axis equal to  $\beta H_y ev_x/c$ .

We see in this way that a magnetic field in one set of axes becomes a magnetic field and an electric field in another set. Magnetic and electric fields are therefore relative quantities, but the electromagnetic field tensor  $t$  is the same in any set of axes, since it is equal to the curl of the Minkowski vector potential, and the latter is a four-dimensional vector.

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## CHAPTER XIX

# General Relativity and Gravitation

### 1. Principle of Equivalence.

We have seen that, since a gravitational field gives equal accelerations to material particles of all kinds, it is impossible, by observations on the motion of such particles in a laboratory, to distinguish between accelerations due to a gravitational field and accelerations due to an acceleration of the laboratory.

Einstein put forward a generalization of this principle which is known as the *principle of equivalence*. According to the principle of equivalence, the effects due to a uniform gravitational field are precisely the same as those due to a uniform acceleration of the material frame of reference relatively to which the phenomena are observed.

The uniform acceleration is supposed to be equal and opposite to the acceleration which the gravitational field gives to a particle of any kind. It follows that if a frame of reference moves with an acceleration equal to, and in the same direction as, that due to the gravitational field, the effects due to the field will be equal and opposite to those due to the acceleration, and there will be no observable effects.

The principle of equivalence is supposed to be true for electrical and optical phenomena as well as for the motion of material particles.

According to this, if the gravitational fields of the sun and moon were uniform over the whole earth, there would be no observable effects on the earth due to these fields. The tides which are observed are attributed to the variation of the fields with the distance from the sun and moon.

A good illustration of the principle of equivalence is the following, due to Einstein. Imagine an observer working in a large completely closed box, and suppose he knows nothing of what goes on outside the box. Let the box be far from all other material bodies so that there is no gravitational field where it is. Now suppose a rope attached to the box and the rope pulled so that the box is made to move with an acceleration. The observer in the box would find that everything in the box when unsupported moved with an equal and opposite acceleration relative to the box. He would conclude probably that

the box was in a gravitational field. To keep a body at rest in the box would require a force proportional to the mass of the body. The observer could measure the acceleration by means of a pendulum, just as the acceleration of gravity is measured on the earth. If he could observe the path of a ray of light with sufficient accuracy, he would find it not exactly straight because of the acceleration of the box. He might conclude that the refractive index of the space in his box varied slightly from point to point.

Suppose now that the box is supported in a uniform gravitational field so that it remains at rest. To the observer in the box, according to the principle of equivalence, everything will be the same as in the first case. It follows, for example, that light rays must be slightly deviated by a gravitational field.

The principle of equivalence does not mean that it is never possible to distinguish between gravitational fields and accelerations of the frame of reference. For example, the acceleration of gravity  $g$  observed on the earth cannot be attributed to an upward acceleration of the earth's surface equal to  $g$ , because such an acceleration would mean that the diameter of the earth was increasing all the time, which is of course impossible.

The principle of equivalence suggests that producing a gravitational field in a space is analogous to changing the frame of reference or co-ordinate system used to describe phenomena in the space. In the four-dimensional Minkowski world of the special theory of relativity the world lines of the particles represent phenomena independently of any co-ordinate system. Let us suppose that the system we are considering consists of particles subject only to gravitational forces—that is, suppose that there are no electric, magnetic, or other forces besides gravitational in the system. Einstein supposes that the special theory of relativity is true for such a system only when the masses of the particles are so small and the distances between them so large that the gravitational forces are negligible. In this case the world lines of the particles are all straight in any system of rectangular co-ordinates  $x, y, z, \tau$  which may be adopted. The interval between two points  $A$  and  $B$  measured along a world line joining the points is equal to  $\int_A^B ds$ , and for a straight line between  $A$  and  $B$   $\delta \int_A^B ds = 0$ , where  $\delta$  denotes a variation from the straight line to any infinitely near line from  $A$  to  $B$ .

A line for which  $\int_A^B ds$  is stationary, or for which  $\delta \int_A^B ds = 0$ , is called a *geodetic line* or *geodesic*.

## 2. Curvilinear Co-ordinates. World Lines and Geodesics.

The interval  $ds$  between two neighbouring points is the same in any system of rectangular co-ordinates, so that this is true also of  $\int_A^B ds$ .

Suppose that we use a system of curvilinear co-ordinates instead of the rectangular ones. Then, since  $ds$  has the same value in any co-ordinate system,  $\int_A^B ds$  along a world line will still be stationary in the curvilinear system. The world line will still be given by  $\delta \int_A^B ds = 0$ , where  $\delta$  denotes a variation due to changing from the world line between  $A$  and  $B$  to any other line between  $A$  and  $B$  which is very near to the world line.

Now according to the principle of equivalence the effects due to a gravitational field are the same as the effects due to changing the co-ordinate system in a suitable way, so that we may assume that in a gravitational field the world line of a particle will be given by  $\delta \int_A^B ds = 0$ .

If, however, we use rectangular co-ordinates in the presence of a gravitational field  $\delta \int_A^B ds = 0$  will give straight lines, whereas the world lines in a gravitational field will not be straight, because the gravitational forces will produce curvature of the lines. We conclude, therefore, that rectangular co-ordinates are impossible in a gravitational field. This means that the gravitational field modifies space in such a way that it is impossible to choose co-ordinates in which  $ds^2$  is everywhere equal to  $dx^2 + dy^2 + dz^2 + d\tau^2$ .

Since a very short element of any curve may be regarded as straight, we see that it is possible to choose rectangular co-ordinates in which any particular element  $ds$  of a world line is given by  $ds^2 = dx^2 + dy^2 + dz^2 + d\tau^2$ , but the same rectangular co-ordinates cannot be used for the successive elements  $ds$  of the line.

To see what this means let us take a simple case from ordinary geometry. Consider two points  $A$  and  $B$  lying in a plane, and let  $ds$  be an element of any line from  $A$  to  $B$ . If  $\delta \int_A^B ds = 0$ , the line is as short as possible. Let  $x, y$  be the rectangular co-ordinates of a point on the plane, so that  $ds^2 = dx^2 + dy^2$ . Suppose we draw two sets of equidistant parallel straight lines on the plane, one set parallel to the  $x$  axis and the other to the  $y$  axis. Let these lines be given by  $x = 0, 1, 2, 3, 4 \dots$  and  $y = 0, 1, 2, 3, 4 \dots$ , and let the lines be numbered with numbers equal to the values of  $x$  or  $y$  at each line.

Suppose now that the sets of lines are distorted in any way so that they become curved. If, for example, they were drawn on a thin

sheet of rubber, by stretching the rubber they could be changed into two sets of curves. The small equal squares between the lines would be changed into unequal parallelograms. We suppose the line between the two points  $A$  and  $B$  to be fixed and not moved in any way by the distortion of the two sets of lines.

We may use the numbers on the curved lines as co-ordinates of a point on the plane if we suppose that the distortion did not move any of the lines out of the plane. Thus a point half-way between the two  $x$  lines numbered 5 and 6 and the two  $y$  lines numbered 9 and 10 would have co-ordinates 5.5 and 9.5. It is clear that any such change from rectangular to curvilinear co-ordinates in the plane makes no difference to  $\int_A^B ds$  and that  $\delta \int_A^B ds = 0$  is still the condition for the line joining  $AB$  to be as short as possible, because the length of  $ds$  is the same in whatever system of co-ordinates it is expressed. If the co-ordinates of  $A$  are  $x_1, y_1$  in the rectangular system and those of  $B$   $x_2, y_2$ , then  $AB^2 = (x_2 - x_1)^2 + (y_2 - y_1)^2$ , and the equation of the straight line  $AB$  is

$$y - y_1 = (y_2 - y_1) \frac{x - x_1}{x_2 - x_1},$$

or  $y = ax + b$ , where  $a$  and  $b$  are constants. In the curvilinear system the equation of the line  $AB$  will not be linear so that the line may be said to be curved relatively to the curvilinear co-ordinates if we regard a line given by a linear equation as straight. The line given by  $\delta \int_A^B ds = 0$  will be straight in rectangular co-ordinates and curved in curvilinear co-ordinates, but in any co-ordinates it will be the shortest line between  $A$  and  $B$ .

Such a change to curvilinear co-ordinates in a plane is analogous to changing from rectangular co-ordinates  $x, y, z, \tau$  to curvilinear co-ordinates in the Minkowski world when there is no gravitational field. In the curvilinear system particles move along curved paths as though acted on by a field of force, or in other words the use of a co-ordinate system which is accelerated produces an apparent field which is exactly like a gravitational field.

Suppose now that when the two sets of equidistant parallel lines in the plane are distorted the plane itself is also distorted into a curved surface. Then a line from  $A$  to  $B$  on the surface cannot be straight, but  $\delta \int_A^B ds = 0$  will still give the shortest line from  $A$  to  $B$  on the surface.

On a curved surface it is not in general possible to set up a system of rectangular co-ordinates in which  $ds^2 = dx^2 + dy^2$  everywhere, but any small element of the surface can be regarded as plane, and a small local set of rectangular axes can be drawn on it in which  $ds^2 = dx^2 + dy^2$ .

According to Einstein's theory, the distortion in this case is analogous to the introduction of a gravitational field into the Minkowski world. Einstein supposes that the gravitational field distorts or curves space so that rectangular co-ordinates in which  $ds^2 = dx^2 + dy^2 + dz^2 + d\tau^2$ , everywhere, become impossible, but  $\delta \int_A^B ds = 0$  still gives the position of the world line of a particle from  $A$  to  $B$ .

According to this theory, then, the world line of a particle between two points  $A$  and  $B$  is always given by  $\delta \int_A^B ds = 0$ .

The gravitational field is a curvature or distortion of the Minkowski four-dimensional world, and the world lines are geodesics in this distorted space. The equation  $\delta \int_A^B ds = 0$  is purely geometrical, and so all particles of whatever mass or nature move in the same way in a given field. That is, the gravitational acceleration is the same for all kinds of matter, as is found experimentally to be the case.

As we have seen, along the path of a ray of light in the Minkowski world, when there is no gravitational field, we have  $ds^2 = dx^2 + dy^2 + dz^2 - c^2 dt^2$ , where  $icdt$  has been put for  $d\tau$ , and since in this case

$$v^2 = \frac{dx^2 + dy^2 + dz^2}{dt^2} = c^2,$$

we get  $ds = 0$  for any element of the line. This result of course must be true in any system of co-ordinates and, according to the principle of equivalence, it must also be true in a gravitational field. The world line of a ray of light is therefore a geodesic of zero length.

To determine the path of a particle or ray of light in a gravitational field it is therefore necessary to find in what way space is distorted near material bodies, so that  $ds$  may be expressed in terms of any set of co-ordinates chosen and the equation of the world line between two points found in terms of the co-ordinates by means of the equation  $\delta \int_A^B ds = 0$ . The co-ordinates used may be any which are possible in the curved space just as, for example, any co-ordinates may be used to fix the positions of points on the surface of a sphere, provided they conform to the geometry of the spherical surface.

### 3. General Expression for $ds^2$ . Einstein's Problem..

In the curved Minkowski world any very small region may be regarded as not distorted, and a small local system of rectangular co-ordinates may be supposed drawn in it. In these local co-ordinates  $ds^2 = dx^2 + dy^2 + dz^2 + d\tau^2$ . Now let  $x_1, x_2, x_3, x_4$  be taken as the co-ordinates in a system which can be used throughout the space. Then in the small region of the co-ordinates  $x, y, z, \tau$  these rectangular

co-ordinates must be functions of  $x_1, x_2, x_3, x_4$ . Hence we have

$$dx = \frac{\partial x}{\partial x_1} dx_1 + \frac{\partial x}{\partial x_2} dx_2 + \frac{\partial x}{\partial x_3} dx_3 + \frac{\partial x}{\partial x_4} dx_4,$$

with similar equations for  $dy, dz$ , and  $d\tau$ .

Substituting these values of  $dx, dy, dz$ , and  $d\tau$  in the equation  $ds^2 = dx^2 + dy^2 + dz^2 + d\tau^2$ , we get

$$\begin{aligned} ds^2 = & g_{11}dx_1^2 + g_{22}dx_2^2 + g_{33}dx_3^2 + g_{44}dx_4^2 \\ & + 2g_{12}dx_1dx_2 + 2g_{13}dx_1dx_3 + 2g_{14}dx_1dx_4 \\ & + 2g_{23}dx_2dx_3 + 2g_{24}dx_2dx_4 + 2g_{34}dx_3dx_4, \end{aligned}$$

where the  $g$ 's are functions of the co-ordinates  $x_1, x_2, x_3, x_4$ . For

example,  $g_{11} = \left(\frac{\partial x}{\partial x_1}\right)^2 + \left(\frac{\partial y}{\partial x_1}\right)^2 + \left(\frac{\partial z}{\partial x_1}\right)^2 + \left(\frac{\partial \tau}{\partial x_1}\right)^2$ , and  $x, y, z, \tau$  are functions of  $x_1, x_2, x_3, x_4$ . This expression for  $ds^2$  may be written  $ds^2 = \Sigma g_{\mu\nu} dx_\mu dx_\nu$ , in which it is understood that all the sixteen combinations of the four values 1, 2, 3, 4 of  $\mu$  and  $\nu$  are to be summed, and that  $g_{\mu\nu} = g_{\nu\mu}$ . An expression for  $ds^2$  of this form holds of course in any system of co-ordinates.

The relations between the  $g$ 's and the properties of the curved space had been previously worked out by Riemann and other mathematicians, so that Einstein was able to make use of their results. The problem is to find the relations between the  $g$ 's which hold in any possible co-ordinate system in a gravitational field. The relations between the  $g$ 's are sets of differential equations, and Einstein selected a set which seemed likely to be the correct set. In making this more or less arbitrary choice he was guided by the knowledge that Newton's law of gravitation is certainly a close approximation to the truth, so that it was necessary to choose equations which gave results differing little from Newton's theory. Also, in the absence of a gravitational field he supposed the four-dimensional space to be undistorted, so that the differential relations between the  $g$ 's had to be such that they reduced to the relations for an undistorted space at great distances from matter.

Einstein succeeded in finding a set of equations satisfying these conditions, and he then worked out the theory of the motion of a small planet round a large attracting mass, and the path of a ray of light near the sun. The results obtained have been found to agree with the facts, and his theory is therefore regarded as probably correct.

#### 4. Theory of Tensors.

Thus Einstein's theory depends on the differential geometry of four-dimensional space, which in turn depends on the theory of tensors. In a space of four dimensions let  $x_1, x_2, x_3, x_4$  be the co-ordinates of a

point, and let the components of a small displacement be  $dx_1, dx_2, dx_3, dx_4$ . If now we change to any other set of co-ordinates  $x'_1, x'_2, x'_3, x'_4$  the components of the small displacement will be given by

$$dx'_1 = \frac{\partial x'_1}{\partial x_1} dx_1 + \frac{\partial x'_1}{\partial x_2} dx_2 + \frac{\partial x'_1}{\partial x_3} dx_3 + \frac{\partial x'_1}{\partial x_4} dx_4,$$

with similar equations for  $dx'_2, dx'_3$ , and  $dx'_4$ . These equations may be written

$$dx'_\mu = \frac{\partial x'_\mu}{\partial x_\sigma} dx_\sigma,$$

where  $\sigma$  is to be given the values 1, 2, 3, 4, and the four terms are to be added.

In what follows it will be understood that, when a suffix occurs twice in a term, the term stands for the sum of its four values corresponding to the four values of the suffix 1, 2, 3, and 4.

A displacement is a vector, and any other vector which is transformed from one set of co-ordinates to another in the same way as a displacement is called a *contravariant vector*. A contravariant vector with components  $A^1, A^2, A^3, A^4$  referred to  $x_1, x_2, x_3, x_4$  transforms into one with components  $A'^1, A'^2, A'^3, A'^4$  referred to  $x'_1, x'_2, x'_3, x'_4$  where

$$A'^1 = \frac{\partial x'_1}{\partial x_1} A^1 + \frac{\partial x'_1}{\partial x_2} A^2 + \frac{\partial x'_1}{\partial x_3} A^3 + \frac{\partial x'_1}{\partial x_4} A^4,$$

with similar equations for  $A'^2, A'^3$ , and  $A'^4$ .

Thus a contravariant vector  $A^\mu$  transforms into

$$A'^\mu = \frac{\partial x'_\mu}{\partial x_\sigma} A^\sigma,$$

where  $\mu = 1, 2, 3$ , or 4 and  $\sigma = 1, 2, 3$ , or 4. Since  $\sigma$  appears twice the term  $\frac{\partial x'_\mu}{\partial x_\sigma} A^\sigma$  is understood to be summed.

If  $\phi$  is a scalar function of position, the vector having components  $\frac{\partial \phi}{\partial x'_1}, \frac{\partial \phi}{\partial x'_2}, \frac{\partial \phi}{\partial x'_3}, \frac{\partial \phi}{\partial x'_4}$  is transformed by the formula

$$\frac{\partial \phi}{\partial x'_1} = \frac{\partial x_1}{\partial x'_1} \frac{\partial \phi}{\partial x_1} + \frac{\partial x_2}{\partial x'_1} \frac{\partial \phi}{\partial x_2} + \frac{\partial x_3}{\partial x'_1} \frac{\partial \phi}{\partial x_3} + \frac{\partial x_4}{\partial x'_1} \frac{\partial \phi}{\partial x_4},$$

since we have  $d\phi = \frac{\partial \phi}{\partial x_1} dx_1 + \frac{\partial \phi}{\partial x_2} dx_2 + \frac{\partial \phi}{\partial x_3} dx_3 + \frac{\partial \phi}{\partial x_4} dx_4$ ; with similar equations for  $\frac{\partial \phi}{\partial x'_2}, \frac{\partial \phi}{\partial x'_3}$ , and  $\frac{\partial \phi}{\partial x'_4}$ . A vector which transforms in this

way is called a *covariant vector*. If  $A_\mu$  denotes a component of a covariant vector, then

$$A'_\mu = \frac{\partial x_\sigma}{\partial x'_\mu} A_\sigma.$$

Contravariant vectors are indicated by a raised suffix and covariant vectors by a lowered suffix. The contravariant vector  $dx_\mu$ , however, is usually written with a lowered suffix.

*Tensors* are quantities having components which transform in the same way as the products of the components of two or more vectors. A tensor which transforms like the products of the components of two covariant vectors is written  $A_{\mu\nu}$ , and transforms thus:

$$A'_{\mu\nu} = \frac{\partial x_\sigma}{\partial x'_\mu} \frac{\partial x_\tau}{\partial x'_\nu} A_{\sigma\tau}.$$

In this expression  $\sigma$  and  $\tau$  occur twice, so that there are sixteen terms to be summed. The covariant tensor  $A_{\mu\nu}$  has sixteen components,

$$\begin{array}{cccc} A_{11} & A_{12} & A_{13} & A_{14} \\ A_{21} & A_{22} & A_{23} & A_{24} \\ A_{31} & A_{32} & A_{33} & A_{34} \\ A_{41} & A_{42} & A_{43} & A_{44}. \end{array}$$

In the same way, we define contravariant tensors  $A^{\mu\nu}$ , which transform like the products of components of contravariant vectors, so that

$$A'^{\mu\nu} = \frac{\partial x'_\mu}{\partial x_\sigma} \frac{\partial x'_\nu}{\partial x_\tau} A^{\sigma\tau};$$

and mixed tensors  $A^\nu_\mu$ , for which

$$A'^\nu_\mu = \frac{\partial x_\sigma}{\partial x'_\mu} \frac{\partial x'_\nu}{\partial x_\tau} A^\tau_\sigma.$$

Thus any component of a tensor in co-ordinates  $x'_1, x'_2, x'_3, x'_4$  is equal to the sum of a number of terms each of which is proportional to one of its components in any other co-ordinates  $x_1, x_2, x_3, x_4$ . It follows that if a tensor is zero in any co-ordinates, that is, if all its components are zero, it will be zero in any other co-ordinate system.

If then any law of nature expressing results, relatively to a material frame of reference, in terms of co-ordinates  $x_1, x_2, x_3, x_4$  can be expressed by an equation  $T = 0$ , where  $T$  is a tensor, then in terms of any other co-ordinates  $x'_1, x'_2, x'_3, x'_4$  the law will be expressed by  $T' = 0$ , where  $T'$  is the tensor into which  $T$  is transformed by the change from  $x_1, x_2, x_3, x_4$  to  $x'_1, x'_2, x'_3, x'_4$ .

According to Einstein's general principle of relativity all the laws of nature can be expressed by equations of the form  $T = 0$ , so that all such laws, when so expressed, are independent of the co-ordinate

system used. In particular the law of gravitation must be capable of being put into the form  $T = 0$  like any other law of nature.

Tensors having two suffixes are said to be of the second rank, and vectors may be called tensors of the first rank. Tensors which transform like the products of more than two vector components may be defined in a similar way.

The product of a component  $A_{\mu\nu}$  of a tensor with a component  $B_{\sigma\tau}$  of another tensor is a component of a tensor ( $A_{\mu\nu}B_{\sigma\tau}$ ), the components of which are all the products of the components like  $A_{\mu\nu}$  with those like  $B_{\sigma\tau}$ . The product of two vectors  $A_\mu$  and  $B_\nu$  is a tensor  $A_\mu B_\nu = C_{\mu\nu}$ . The product of  $A_{\mu\nu}$  and  $B_\lambda^\rho$  is  $A_{\mu\nu}B_\lambda^\rho = C_{\mu\nu\lambda}^\rho$ .

If we multiply  $A_\mu$  by  $B^\mu$  we get  $A_\mu B^\mu$ , which is equal to

$$A_1B^1 + A_2B^2 + A_3B^3 + A_4B^4,$$

and so is a scalar quantity having only one component or value. The product  $A_\mu B^\mu$  is called the *inner product*, to distinguish it from the ordinary product  $A_\mu B^\nu$ , which is a tensor having sixteen components, viz.:

$$\begin{array}{cccc} A_1B^1 & A_1B^2 & A_1B^3 & A_1B^4 \\ A_2B^1 & A_2B^2 & A_2B^3 & A_2B^4 \\ A_3B^1 & A_3B^2 & A_3B^3 & A_3B^4 \\ A_4B^1 & A_4B^2 & A_4B^3 & A_4B^4. \end{array}$$

If an upper and lower suffix of a mixed tensor are both denoted by the same letter then the summation rule makes the tensor equal to the sum of four tensors; thus, for example, if in  $A_{\mu\nu}^\sigma$  we put  $\sigma = \nu$  we get

$$A_{\mu\nu}^\nu = A_{\mu 1}^1 + A_{\mu 2}^2 + A_{\mu 3}^3 + A_{\mu 4}^4.$$

The sum may be denoted by  $A_\mu$ , and so denotes a vector with components  $A_1, A_2, A_3, A_4$ . Thus putting  $\sigma = \nu$  in  $A_{\mu\nu}^\sigma$  changes it to a vector  $A_\mu$ . This operation is called *contraction*. If we contract  $A_\mu^\nu$  we get

$$A_1^1 + A_2^2 + A_3^3 + A_4^4,$$

which is an *invariant* (that is, retains the same value in all systems of co-ordinates), because

$$A_\mu'^\mu = \frac{\partial x_\sigma}{\partial x'_\mu} \frac{\partial x'_\mu}{\partial x_\sigma} A_\sigma^\sigma = A_\sigma^\sigma.$$

It is important to be able to find out whether any quantity is a tensor or not. This can be done by finding its equations of transformation, or by expressing it as the sum or product of quantities known to be tensors. A quantity which on inner multiplication by any contravariant or covariant vector always gives a tensor is a tensor.

Thus let the product  $A_{\mu\nu}B^\nu$  be a covariant vector for any  $B^\nu$ . Then

$$A'_{\mu\nu}B'^\nu = \frac{\partial x_\sigma}{\partial x'_\mu} A_{\sigma\tau} B^\tau.$$

Also

$$B^\tau = \frac{\partial x_\tau}{\partial x'_\nu} B'^\nu,$$

so that

$$B'^\nu (A'_{\mu\nu} - \frac{\partial x_\sigma}{\partial x'_\mu} \frac{\partial x_\tau}{\partial x'_\nu} A_{\sigma\tau}) = 0.$$

Since  $B'^\nu$  is arbitrary, it follows that

$$A'_{\mu\nu} = \frac{\partial x_\sigma}{\partial x'_\mu} \frac{\partial x_\tau}{\partial x'_\nu} A_{\sigma\tau},$$

which shows that  $A_{\mu\nu}$  is a tensor.

### 5. The Fundamental Tensor.

Consider the equation  $ds^2 = g_{\mu\nu} dx_\mu dx_\nu$ , where  $g_{\mu\nu} = g_{\nu\mu}$ . It can be shown at once by actual transformation that  $g_{\mu\nu}$  is a covariant tensor. Thus, since

$$dx_\mu = \frac{\partial x_\mu}{\partial x'_\sigma} dx'_\sigma, \text{ and } dx_\nu = \frac{\partial x_\nu}{\partial x'_\tau} dx'_\tau,$$

we have

$$g_{\mu\nu} dx_\mu dx_\nu = g_{\mu\nu} \frac{\partial x_\mu}{\partial x'_\sigma} \frac{\partial x_\nu}{\partial x'_\tau} dx'_\sigma dx'_\tau.$$

If this is written in the form  $g'_{\sigma\tau} dx'_\sigma dx'_\tau$ , we therefore have

$$g'_{\sigma\tau} = \frac{\partial x_\mu}{\partial x'_\sigma} \frac{\partial x_\nu}{\partial x'_\tau} g_{\mu\nu},$$

which shows that  $g_{\mu\nu}$  is a covariant tensor. It is called the fundamental tensor. The product  $g_{\mu\nu} dx_\mu dx_\nu$  is thus of the form  $A_{\mu\nu} B^\mu C^\nu$  and so is an inner product, and invariant, as it should be since  $ds$  is the same in any co-ordinates.

The determinant

$$\begin{vmatrix} g_{11} & g_{12} & g_{13} & g_{14} \\ g_{21} & g_{22} & g_{23} & g_{24} \\ g_{31} & g_{32} & g_{33} & g_{34} \\ g_{41} & g_{42} & g_{43} & g_{44} \end{vmatrix} = |g_{\mu\nu}|$$

is denoted by  $g$ . If we erase from this determinant the  $i$ th row and the  $k$ th column, and multiply by  $(-1)^{i+k}$ , we get a minor  $D_{ik}$  of the determinant. This minor divided by the determinant is denoted by  $g^{ik}$ , so that  $g^{ik} = D_{ik}/g$ . Thus, for example,

$$g^{23} = - \begin{vmatrix} g_{11} & g_{12} & g_{14} \\ g_{31} & g_{32} & g_{34} \\ g_{41} & g_{42} & g_{44} \end{vmatrix} \div |g_{\mu\nu}|.$$

From this it follows that  $g_{\sigma}^{\nu} = g_{\mu\sigma} g^{\mu\nu} = 1$  when  $\sigma = \nu$ , because

$$g_{\mu\sigma} g^{\mu\nu} = g_{1\sigma} g^{1\nu} + g_{2\sigma} g^{2\nu} + g_{3\sigma} g^{3\nu} + g_{4\sigma} g^{4\nu}$$

and

$$g = \sum_{k=1}^{k=4} g_{ik} D_{ik} = \sum_{i=1}^{i=4} g_{ik} D_{ik}.$$

In the same way  $g_{\mu\sigma} g^{\mu\nu} = 0$ , i.e.  $g_{\sigma}^{\nu} = 0$ , when  $\sigma$  is not equal to  $\nu$ , because  $g_{\mu\sigma} D_{\mu\nu}$  gives a determinant with two columns identical, which is therefore equal to zero.

Hence, if  $A^{\nu}$  is any contravariant vector,

$$g_{\sigma}^{\nu} A^{\sigma} = g_1^{\nu} A^1 + g_2^{\nu} A^2 + g_3^{\nu} A^3 + g_4^{\nu} A^4 = A^{\nu},$$

since  $g_{\sigma}^{\nu} = 0$  when  $\sigma \neq \nu$ , and  $g_{\sigma}^{\nu} = 1$  when  $\sigma = \nu$ .

The three fundamental tensors are

$$g_{\mu\nu}, g_{\mu}^{\nu}, g^{\mu\nu}.$$

From any covariant tensor  $A_{\mu\nu}$  we can get a mixed tensor  $A^{\nu}_{\mu} = g^{\nu\alpha} A_{\alpha\mu}$ , a contravariant tensor  $A^{\mu\nu} = g^{\mu\alpha} g^{\nu\beta} A_{\alpha\beta}$ , and an invariant or scalar  $A = g^{\mu\nu} A_{\mu\nu}$ .

## 6. Equations of a Geodesic. Christoffel's Symbols.

The world line of a particle is a geodesic determined by  $\delta \int_A^B ds = 0$ , where  $\delta$  indicates a variation from the line itself between two points  $A$  and  $B$  on it to any other line which is very close to the world line. The variations are zero at  $A$  and  $B$ .

We have  $ds^2 = g_{\mu\nu} dx_{\mu} dx_{\nu} = g_{hk} dx_h dx_k$ , so that  $1 = g_{hk} \frac{dx_h}{ds} \frac{dx_k}{ds}$ .

The geodesics are therefore determined by the equation

$$0 = \delta \int_A^B g_{hk} \frac{dx_h}{ds} \frac{dx_k}{ds} ds,$$

or

$$0 = \int_A^B \delta \left( g_{hk} \frac{dx_h}{ds} \frac{dx_k}{ds} \right) ds.$$

Now

$$\delta \left( g_{hk} \frac{dx_h}{ds} \frac{dx_k}{ds} \right) = \delta g_{hk} \frac{dx_h}{ds} \frac{dx_k}{ds} + g_{hk} \left\{ \frac{dx_h}{ds} \delta \left( \frac{dx_k}{ds} \right) + \frac{dx_k}{ds} \delta \left( \frac{dx_h}{ds} \right) \right\}.$$

Also  $\delta g_{hk} = \frac{\partial g_{hk}}{\partial x_p} \delta x_p$ , and the last two terms in the above equation are equal (as is seen at once by simply interchanging  $h$  and  $k$ , which leaves the same summations as before), so that

$$\delta \left( g_{hk} \frac{dx_h}{ds} \frac{dx_k}{ds} \right) = \frac{\partial g_{hk}}{\partial x_p} \delta x_p \frac{dx_h}{ds} \frac{dx_k}{ds} + 2g_{hp} \frac{dx_h}{ds} \delta \left( \frac{dx_p}{ds} \right).$$

In the last term,  $p$  has been written for convenience instead of  $k$ , but this obviously makes no difference. But

$$\delta \left( \frac{dx_p}{ds} \right) = \frac{d}{ds} (\delta x_p),$$

so that on multiplying by  $ds$  and integrating between  $A$  and  $B$  we get, for the geodesics,

$$\int_A^B \frac{\partial g_{hk}}{\partial x_p} \frac{dx_h}{ds} \frac{dx_k}{ds} \delta x_p ds + 2 \int_A^B g_{hp} \frac{dx_h}{ds} \frac{d}{ds} (\delta x_p) ds = 0.$$

Integrating the second integral by parts, and remembering that the variation  $\delta$  is zero at  $A$  and  $B$ , we get

$$\int_A^B \left\{ \frac{\partial g_{hk}}{\partial x_p} \frac{dx_h}{ds} \frac{dx_k}{ds} - 2 \frac{d}{ds} \left( g_{hp} \frac{dx_h}{ds} \right) \right\} \delta x_p ds = 0.$$

Since the variations  $\delta x_p$  are arbitrary, it follows that

$$\frac{\partial g_{hk}}{\partial x_p} \frac{dx_h}{ds} \frac{dx_k}{ds} - 2 \frac{d}{ds} \left( g_{hp} \frac{dx_h}{ds} \right) = 0,$$

or 
$$g_{hp} \frac{d^2 x_h}{ds^2} + \frac{dg_{hp}}{ds} \frac{dx_h}{ds} - \frac{1}{2} \frac{dx_h}{ds} \frac{dx_k}{ds} \frac{\partial g_{hk}}{\partial x_p} = 0.$$

But 
$$\frac{dg_{hp}}{ds} = \frac{\partial g_{hp}}{\partial x_k} \frac{dx_k}{ds},$$

and 
$$\frac{\partial g_{hp}}{\partial x_k} \frac{dx_k}{ds} \frac{dx_h}{ds} = \frac{\partial g_{kp}}{\partial x_h} \frac{\partial x_h}{\partial s} \frac{\partial x_k}{\partial s},$$

so that, putting 
$$\left[ \begin{smallmatrix} h & k \\ p \end{smallmatrix} \right] = \frac{1}{2} \left( \frac{\partial g_{hp}}{\partial x_k} + \frac{\partial g_{kp}}{\partial x_h} - \frac{\partial g_{hk}}{\partial x_p} \right),$$

we get 
$$g_{hp} \frac{d^2 x_h}{ds^2} + \left[ \begin{smallmatrix} h & k \\ p \end{smallmatrix} \right] \frac{dx_h}{ds} \frac{dx_k}{ds} = 0.$$

The symbol  $\left[ \begin{smallmatrix} h & k \\ p \end{smallmatrix} \right]$  is called a *Christoffel's three-index symbol*. Another

kind of three-index symbol is defined by

$$\left\{ \begin{smallmatrix} h & k \\ r \end{smallmatrix} \right\} = g^{rp} \left[ \begin{smallmatrix} h & k \\ p \end{smallmatrix} \right].$$

If now we multiply the equation

$$g_{hp} \frac{d^2 x_h}{ds^2} + \left[ \begin{smallmatrix} h & k \\ p \end{smallmatrix} \right] \frac{dx_h}{ds} \frac{dx_k}{ds} = 0$$

by  $g^{rp}$ , and sum for  $p$  in the usual way, we get, since  $g_h^r = g_{hp} g^{rp}$  is equal to unity when  $r = h$  and zero when  $r \neq h$ ,

$$\frac{d^2 x_r}{ds^2} = - \left\{ \begin{matrix} h & k \\ r \end{matrix} \right\} \frac{dx_h}{ds} \frac{dx_k}{ds}.$$

There are four equations of this type, corresponding to the values 1, 2, 3, 4 of  $r$ . Since  $ds$  is an element of a world line, these equations may be said to be the equations of motion of a particle moving along the world line of which  $ds$  is an element. When the  $g$ 's are constants,  $\left\{ \begin{matrix} h & k \\ r \end{matrix} \right\} = 0$ , so that  $\frac{d^2 x_r}{ds^2} = 0$ , which indicates that the world lines are straight, or that the particle moves with uniform velocity in a straight line.

## 7. Covariant Differentiation.

Let  $\phi$  be an invariant function of position so that, since  $ds$  is also invariant,  $d\phi/ds$  is invariant.  $d\phi/ds$  is a vector and its components are  $\partial\phi/\partial x_1$ ,  $\partial\phi/\partial x_2$ ,  $\partial\phi/\partial x_3$ ,  $\partial\phi/\partial x_4$ . The components of course depend on the co-ordinates chosen, but  $d\phi/ds$  is the same in any co-ordinates.

We have 
$$\frac{d\phi}{ds} = \frac{\partial\phi}{\partial x_h} \frac{dx_h}{ds}.$$

Let  $\psi = d\phi/ds$ , so that

$$\frac{d^2\phi}{ds^2} = \frac{d\psi}{ds} = \frac{\partial\psi}{\partial x_k} \frac{dx_k}{ds}.$$

Also 
$$\frac{\partial\psi}{\partial x_k} = \frac{\partial}{\partial x_k} \left( \frac{d\phi}{ds} \right) = \frac{\partial^2\phi}{\partial x_h \partial x_k} \frac{dx_h}{ds} + \frac{\partial\phi}{\partial x_h} \frac{\partial}{\partial x_k} \left( \frac{dx_h}{ds} \right).$$

Hence 
$$\frac{d^2\phi}{ds^2} = \frac{\partial^2\phi}{\partial x_h \partial x_k} \frac{\partial x_h}{ds} \frac{dx_k}{ds} + \frac{\partial\phi}{\partial x_h} \frac{d^2 x_h}{ds^2}.$$

In the last term, of course,  $h$  can be changed to  $r$  if desired. Using the equation

$$\frac{d^2 x_r}{ds^2} = - \left\{ \begin{matrix} h & k \\ r \end{matrix} \right\} \frac{dx_h}{ds} \frac{dx_k}{ds},$$

we then get 
$$\frac{d^2\phi}{ds^2} = \left[ \frac{\partial^2\phi}{\partial x_h \partial x_k} - \left\{ \begin{matrix} h & k \\ r \end{matrix} \right\} \frac{\partial\phi}{\partial x_r} \right] \frac{dx_h}{ds} \frac{dx_k}{ds}.$$

Now  $d^2\phi/ds^2$  is an invariant or tensor of zero rank, and  $dx_h$  and  $dx_k$  are arbitrary contravariant vectors, so that  $\frac{\partial^2\phi}{\partial x_h \partial x_k} - \left\{ \begin{matrix} h & k \\ r \end{matrix} \right\} \frac{\partial\phi}{\partial x_r}$  must be a covariant tensor of the second rank (cf. the end of section 4). If

then we put  $A_h = \partial\phi/\partial x_h$ , and denote the tensor just obtained by  $A_{hk}$ , we get

$$A_{hk} = \frac{\partial A_h}{\partial x_k} - \left\{ \begin{matrix} h & k \\ & r \end{matrix} \right\} A_r.$$

$A_{hk}$  is called the *covariant derivative* of  $A_h$  with respect to  $x_k$ . When the  $g$ 's are constants so that  $\left\{ \begin{matrix} h & k \\ & r \end{matrix} \right\} = 0$ , then  $A_{hk} = \frac{\partial A_h}{\partial x_k}$ , as in ordinary differentiation.

If  $A_\lambda$ ,  $B_\mu$  are covariant vectors, and  $A_{\lambda\nu}$ ,  $B_{\mu\nu}$  their covariant derivatives, the above formula gives

$$A_{\lambda\nu} B_\mu + A_\lambda B_{\mu\nu} = \frac{\partial(A_\lambda B_\mu)}{\partial x_\nu} - \left\{ \begin{matrix} \lambda & \nu \\ & \epsilon \end{matrix} \right\} A_\epsilon B_\mu - \left\{ \begin{matrix} \mu & \nu \\ & \epsilon \end{matrix} \right\} A_\lambda B_\epsilon.$$

Putting  $A_{\lambda\mu\nu} = A_{\lambda\nu} B_\mu + A_\lambda B_{\mu\nu}$ ,  $A_{\epsilon\mu} = A_\epsilon B_\mu$ , and  $A_{\lambda\epsilon} = A_\lambda B_\epsilon$ ,

we get 
$$A_{\lambda\mu\nu} = \frac{\partial A_{\lambda\mu}}{\partial x_\nu} - \left\{ \begin{matrix} \lambda & \nu \\ & \epsilon \end{matrix} \right\} A_{\epsilon\mu} - \left\{ \begin{matrix} \mu & \nu \\ & \epsilon \end{matrix} \right\} A_{\lambda\epsilon}.$$

This gives the covariant derivative with respect to  $x_\nu$  of the tensor  $A_{\lambda\mu}$ .

## 8. The Riemann Tensor.

Now obtain the second covariant derivative of a vector  $A_\mu$  with respect first to  $x_\nu$ , then to  $x_\sigma$ . Denoting it by  $A_{\mu\nu\sigma}$ , we get

$$A_{\mu\nu\sigma} = \frac{\partial}{\partial x_\sigma} \left[ \frac{\partial A_\mu}{\partial x_\nu} - \left\{ \begin{matrix} \mu & \nu \\ & \rho \end{matrix} \right\} A_\rho \right] - \left\{ \begin{matrix} \mu & \sigma \\ & \epsilon \end{matrix} \right\} \left[ \frac{\partial A_\epsilon}{\partial x_\nu} - \left\{ \begin{matrix} \epsilon & \nu \\ & \rho \end{matrix} \right\} A_\rho \right] \\ - \left\{ \begin{matrix} \nu & \sigma \\ & \epsilon \end{matrix} \right\} \left[ \frac{\partial A_\mu}{\partial x_\epsilon} - \left\{ \begin{matrix} \mu & \epsilon \\ & \rho \end{matrix} \right\} A_\rho \right].$$

Reversing the order of the differentiation, we get a similar expression for  $A_{\mu\sigma\nu}$ , and so obtain

$$A_{\mu\nu\sigma} - A_{\mu\sigma\nu} = A_\rho \left[ \left\{ \begin{matrix} \mu & \sigma \\ & \epsilon \end{matrix} \right\} \left\{ \begin{matrix} \epsilon & \nu \\ & \rho \end{matrix} \right\} - \left\{ \begin{matrix} \mu & \nu \\ & \epsilon \end{matrix} \right\} \left\{ \begin{matrix} \epsilon & \sigma \\ & \rho \end{matrix} \right\} + \frac{\partial}{\partial x_\nu} \left\{ \begin{matrix} \mu & \sigma \\ & \rho \end{matrix} \right\} - \frac{\partial}{\partial x_\sigma} \left\{ \begin{matrix} \mu & \nu \\ & \rho \end{matrix} \right\} \right].$$

The difference  $A_{\mu\nu\sigma} - A_{\mu\sigma\nu}$  is a tensor, and  $A_\rho$  is an arbitrary vector so that we see that the expression in the bracket must be a tensor. It is called the *Riemann tensor*, and is denoted by  $R^\rho_{\mu\nu\sigma}$ , since when multiplied by  $A_\rho$  it gives  $A_{\mu\nu\sigma} - A_{\mu\sigma\nu}$ .

If the  $g$ 's are constants, then all the three index symbols vanish, so that the Riemann tensor is then zero. The equations  $R^\rho_{\mu\nu\sigma} = 0$  are a set of differential equations between the  $g$ 's, and since  $R^\rho_{\mu\nu\sigma}$  is a tensor it follows that if the equations  $R^\rho_{\mu\nu\sigma} = 0$  are true in any one set of co-ordinates then they are true in any other set into which the

first set can be transformed. When the  $g$ 's are constants there is no gravitational field, so that the equations  $R_{\mu\nu\sigma}^{\rho} = 0$  are the relations between the  $g$ 's which hold when there is no gravitational field, or when the space is undistorted, or, as we may say, is flat. Of course, if we use curvilinear co-ordinates in flat space, particles will move along curved lines as though acted on by gravitational forces, but these apparent forces are due to the acceleration of the co-ordinates and not to gravitation. If  $R_{\mu\nu\sigma}^{\rho} = 0$  then rectangular axes for which  $ds^2 = dx_1^2 + dx_2^2 + dx_3^2 + dx_4^2$  are possible, since the space is flat.

## 9. Einstein's Law of Gravitation.

We are now in a position to consider Einstein's law of gravitation, or the relations between the  $g$ 's which hold in curved space. The  $g$ 's are analogous to the gravitational potential of Newton's theory. To see this, consider an element  $ds$  of a world line and suppose that its components  $dx_1, dx_2, dx_3, dx_4$  are all zero except  $dx_4$ , so that  $ds^2 = g_{44} dx_4^2$ . Then, if  $dx_4 = icdt$ , we have  $\left(\frac{ds}{dt}\right)^2 = -g_{44} c^2$ . Now if  $\phi$  is the gravitational potential in Newton's theory,  $\frac{1}{2}v^2 = A - \phi$ , where  $v$  is the velocity of a particle of unit mass, and  $A$  is a constant. Hence, putting  $v = ds/dt$ , we get  $\left(\frac{ds}{dt}\right)^2 = 2(A - \phi)$ . Thus we see that  $g_{44}$  is analogous to  $\phi$ . In Newton's theory in empty space we have

$$\frac{\partial^2 \phi}{\partial x^2} + \frac{\partial^2 \phi}{\partial y^2} + \frac{\partial^2 \phi}{\partial z^2} = 0,$$

so that, since Newton's theory is certainly a good approximation to the truth, we should expect the relations between the  $g$ 's in empty space to be differential equations analogous to  $\Delta\phi = 0$ .

Also the relations must be consistent with  $R_{\mu\nu\sigma}^{\rho} = 0$ , but they must be less stringent, so as to admit gravitational fields or distortion of space.

It occurred to Einstein to try the contracted Riemann tensor  $R_{\mu\nu} = 0$ , and this guess has proved satisfactory.  $R_{\mu\nu} = 0$  is a set of differential equations between the  $g$ 's, of the second order like  $\Delta\phi = 0$ , and clearly consistent with  $R_{\mu\nu\sigma}^{\rho} = 0$ . The equations are tensor equations, and therefore consistent with the general principle of relativity. No other tensor has been discovered which can be used instead of  $R_{\mu\nu}$ .

Putting  $\sigma = \rho$  in the expression for  $R_{\mu\nu\sigma}^{\rho}$ , we get

$$R_{\mu\nu} = \frac{\partial}{\partial x_{\nu}} \left\{ \begin{matrix} \mu \rho \\ \rho \end{matrix} \right\} - \frac{\partial}{\partial x_{\rho}} \left\{ \begin{matrix} \mu \nu \\ \rho \end{matrix} \right\} + \left\{ \begin{matrix} \mu \rho \\ \epsilon \end{matrix} \right\} \left\{ \begin{matrix} \nu \epsilon \\ \rho \end{matrix} \right\} - \left\{ \begin{matrix} \mu \nu \\ \epsilon \end{matrix} \right\} \left\{ \begin{matrix} \epsilon \rho \\ \rho \end{matrix} \right\} = 0.$$

According to Einstein, these are the relations between the  $g$ 's which must hold in any case whether the space is curved or flat. In flat space the more stringent condition  $R_{\mu\nu\sigma}^{\rho} = 0$  is true.

The principle of equivalence can now be stated more precisely. The relations  $R_{\mu\nu\sigma}^{\rho} = 0$  and  $R_{\mu\nu} = 0$  are relations between the *second* differential coefficients of the  $g$ 's. It follows that all laws which hold in flat space and depend on the  $g$ 's and their *first* derivatives only will also hold in curved space. Laws which involve the second derivatives of the  $g$ 's need not hold in both flat space and curved space.

### 10. The Field due to a Particle.

Now consider the gravitational field in the space surrounding a single heavy particle. The problem is to find a set of values of the  $g$ 's which satisfy  $R_{\mu\nu} = 0$  and the special conditions of the case.

In flat space we could suppose the particle at the origin and use polar co-ordinates  $r, \theta, \phi, t$ ; that is, let  $x_1 = r, x_2 = \theta, x_3 = \phi, x_4 = ict$ , so that  $ds^2 = dr^2 + r^2 d\theta^2 + r^2 \sin^2 \theta d\phi^2 - c^2 dt^2$ . In this case,  $g_{11} = 1, g_{22} = r^2, g_{33} = r^2 \sin^2 \theta, g_{44} = -c^2$ , and  $g_{\sigma\tau} = 0$  when  $\sigma \neq \tau$ . These values, however, would satisfy  $R_{\mu\nu\sigma}^{\rho} = 0$ , and not merely  $R_{\mu\nu} = 0$ . Let us put  $c = 1$ , and assume (changing for convenience the sign of  $ds^2$ )  $g_{11} = -\epsilon^\lambda, g_{22} = -r^2, g_{33} = -r^2 \sin^2 \theta, g_{44} = \epsilon^\nu$ , and  $g_{\sigma\tau} = 0$  when  $\sigma \neq \tau$ . If these are possible values of the  $g$ 's, they will satisfy the equations  $R_{\mu\nu} = 0$ . It is assumed that  $\lambda$  and  $\nu$  are functions of  $r$  only.

With these  $g$ 's it is easy to calculate the three index symbols. The determinant  $g$  is equal to  $-\epsilon^{\lambda+\nu} r^4 \sin^2 \theta$ , and  $g^{\sigma\sigma} = \frac{1}{g_{\sigma\sigma}}$ , hence we get

$$\left\{ \begin{matrix} \sigma\tau \\ \alpha \end{matrix} \right\} = \frac{1}{2g_{\alpha\alpha}} \left\{ \frac{\partial g_{\alpha\sigma}}{\partial x_\tau} + \frac{\partial g_{\alpha\tau}}{\partial x_\sigma} - \frac{\partial g_{\sigma\tau}}{\partial x_\alpha} \right\},$$

which is not to be summed.

Using this expression we find the following values of  $\left\{ \begin{matrix} \sigma\tau \\ \alpha \end{matrix} \right\}$ :

$$\begin{aligned} \left\{ \begin{matrix} 11 \\ 1 \end{matrix} \right\} &= \frac{1}{2} \frac{d\lambda}{dr}, & \left\{ \begin{matrix} 13 \\ 3 \end{matrix} \right\} &= \frac{1}{r}, \\ \left\{ \begin{matrix} 12 \\ 2 \end{matrix} \right\} &= \frac{1}{r}, & \left\{ \begin{matrix} 14 \\ 4 \end{matrix} \right\} &= \frac{1}{2} \frac{d\nu}{dr}, \\ \left\{ \begin{matrix} 22 \\ 1 \end{matrix} \right\} &= -r\epsilon^{-\lambda}, & \left\{ \begin{matrix} 23 \\ 3 \end{matrix} \right\} &= \cot \theta, \\ \left\{ \begin{matrix} 33 \\ 1 \end{matrix} \right\} &= -r \sin^2 \theta \epsilon^{-\lambda}, & \left\{ \begin{matrix} 33 \\ 2 \end{matrix} \right\} &= -\sin \theta \cos \theta, \\ \left\{ \begin{matrix} 44 \\ 1 \end{matrix} \right\} &= \frac{1}{2} \epsilon^{\nu-\lambda} \frac{d\nu}{dr}. \end{aligned}$$

The rest of the three-index symbols are zero. Using these values of

$\left\{ \begin{smallmatrix} \sigma \tau \\ \alpha \end{smallmatrix} \right\}$  we now find the values of  $R_{\mu\nu}$ . It is found that they are all except  $R_{11}$ ,  $R_{22}$ ,  $R_{33}$ , and  $R_{44}$ .

The values of these are found to be

$$\begin{aligned} R_{11} &= \frac{1}{2} \frac{d^2\nu}{dr^2} - \frac{1}{4} \frac{d\lambda}{dr} \frac{d\nu}{dr} + \frac{1}{4} \left( \frac{d\nu}{dr} \right)^2 - \frac{1}{r} \frac{d\lambda}{dr}, \\ R_{22} &= \epsilon^{-\lambda} \left\{ 1 + \frac{1}{2} r \left( \frac{d\nu}{dr} - \frac{d\lambda}{dr} \right) \right\} - 1, \\ R_{33} &= \sin^2\theta \cdot \epsilon^{-\lambda} \left\{ 1 + \frac{1}{2} r \left( \frac{d\nu}{dr} - \frac{d\lambda}{dr} \right) \right\} - \sin^2\theta, \\ R_{44} &= \epsilon^{\nu-\lambda} \left\{ -\frac{1}{2} \frac{d^2\nu}{dr^2} + \frac{1}{4} \frac{d\lambda}{dr} \frac{d\nu}{dr} - \frac{1}{4} \left( \frac{d\nu}{dr} \right)^2 - \frac{1}{r} \frac{d\nu}{dr} \right\}. \end{aligned}$$

Since each of these is to vanish, we obtain four equations. The first and last equations give  $\frac{\partial\lambda}{\partial r} = -\frac{\partial\nu}{\partial r}$ ; also, when  $r = \infty$ ,  $\lambda$  and  $\nu$  both be zero; therefore  $\lambda = -\nu$ .

The other two equations then give

$$\epsilon^{\nu} \left( 1 + r \frac{\partial\nu}{\partial r} \right) = 1,$$

so that  $\epsilon^{\nu} = 1 - \frac{2m}{r}$ , where  $m$  is an integration constant. This solution satisfies all the equations.

Thus it appears that the forms assumed for the  $g$ 's can satisfy the equations  $R_{\mu\nu} = 0$ , and they give a space symmetrical about the origin, so that they are a possible set for the case of a heavy particle at the origin. The result obtained is

$$ds^2 = - \left( 1 - \frac{2m}{r} \right)^{-1} dr^2 - r^2 d\theta^2 - r^2 \sin^2\theta d\phi^2 + \left( 1 - \frac{2m}{r} \right) dt^2.$$

## 11. Planetary Orbit.

Now suppose that an infinitesimal particle is moving in the space near the heavy particle at the origin. To find its motion we may either use  $\int ds = 0$  or the equations of motion

$$\frac{d^2 x_r}{ds^2} = - \left\{ \begin{smallmatrix} h & k \\ r \end{smallmatrix} \right\} \frac{dx_h}{ds} \frac{dx_k}{ds}.$$

Substituting the values found above for  $\left\{ \begin{smallmatrix} h & k \\ r \end{smallmatrix} \right\}$  we get, when  $r$  so that  $x_r = \theta$ ,

$$\frac{d^2\theta}{ds^2} = \cos\theta \sin\theta \left( \frac{d\phi}{ds} \right)^2 - \frac{2}{r} \frac{dr}{ds} \frac{d\theta}{ds}.$$

If initially the particle is moving in the plane  $\theta = \pi/2$ , then  $d\theta/ds = 0$ , and  $\cos\theta = 0$ , so that initially

$$\frac{d^2\theta}{ds^2} = 0,$$

The particle therefore remains in the plane  $\theta = \pi/2$ . With  $\theta = \pi/2$ , the equations for  $x_3$  and  $x_4$  are

$$\frac{d^2\phi}{ds^2} + \frac{2}{r} \frac{dr}{ds} \frac{d\phi}{ds} = 0,$$

$$\frac{d^2t}{ds^2} + \frac{dv}{dr} \frac{dr}{ds} \frac{dt}{ds} = 0.$$

Integrating these equations we get  $r^2 \frac{d\phi}{ds} = h$ , and  $\frac{dt}{ds} = \frac{a}{\gamma}$ , where  $h$  and  $a$  are constants, and  $\gamma = \epsilon^v = 1 - \frac{2m}{r}$ . The equation for  $ds^2$ , with  $d\theta = 0$  and  $\sin\theta = 1$ , gives  $ds^2 = -\gamma^{-1} dr^2 - r^2 d\phi^2 + \gamma dt^2$ . This with  $\frac{dt}{ds} = \frac{a}{\gamma}$  becomes

$$1 = -\gamma^{-1} \left(\frac{dr}{ds}\right)^2 - r^2 \left(\frac{d\phi}{ds}\right)^2 + a^2 \gamma^{-1}.$$

Since  $\frac{dr}{ds} = \frac{dr}{d\phi} \frac{d\phi}{ds}$ ,  $\frac{d\phi}{ds} = \frac{h}{r^2}$ , and  $\gamma = 1 - \frac{2m}{r}$ , we get

$$\left(\frac{dr}{d\phi}\right)^2 \frac{h^2}{r^4} = a^2 - 1 + \frac{2m}{r} - \frac{h^2}{r^2} + \frac{2mh^2}{r^3}.$$

Putting  $u = r^{-1}$ , this becomes

$$\left(\frac{du}{d\phi}\right)^2 + u^2 = \frac{a^2 - 1}{h^2} + \frac{2mu}{h^2} + 2mu^3.$$

The corresponding equation for Newton's law of gravitation is

$$\left(\frac{du}{d\phi}\right)^2 + u^2 = \frac{a^2 - 1}{h^2} + \frac{2mu}{h^2},$$

where  $m$  is the mass of the particle at the origin, and  $h = \frac{r^2 d\phi}{dt}$ . We

may conclude that  $m$  is the mass of the particle in Einstein's theory. The additional term  $2mu^3$  is very small in the case of the planets of the solar system, and makes no appreciable difference except in the case of the planet Mercury.

On differentiating Einstein's equation with respect to  $\phi$  and dividing by  $2du/d\phi$ , we find

$$\frac{d^2u}{d\phi^2} + u = \frac{m}{h^2} + 3mu^2.$$

The corresponding Newtonian equation is

$$\frac{d^2u}{d\phi^2} + u = \frac{m}{h^2},$$

the solution of which is

$$u = \frac{m}{h^2} (1 + e \cos \phi),$$

representing (if  $e < 1$ ) an ellipse with the attracting particle at one focus, and perihelion at  $\phi = 0$ . The additional term  $3mu^2$  in Einstein's equation is very small in comparison with  $m/h^2$  in planetary orbits. We shall show that it corresponds to a slow rotation of the major axis of the ellipse.

Following the method of successive approximation, we substitute the approximate value of  $u$ , as derived from the Newtonian equation, in Einstein's equation, and so find

$$\frac{d^2u}{d\phi^2} + u = \frac{m}{h^2} + \frac{3m^3}{h^4} (1 + \frac{1}{2}e^2 + 2e \cos \phi + \frac{1}{2}e^2 \cos 2\phi),$$

the solution of which is

$$u = \left\{ \frac{m}{h^2} + \frac{3m^3}{h^4} (1 + \frac{1}{2}e^2) \right\} (1 + e \cos \phi) + \frac{3m^3e}{h^4} (\phi \sin \phi - \frac{1}{6}e \cos 2\phi).$$

At perihelion  $u$  is a maximum, and  $\frac{du}{d\phi} = 0$ . Hence

$$0 = - \left\{ \frac{m}{h^2} + \frac{3m^3}{h^4} (1 + \frac{1}{2}e^2) \right\} e \sin \phi + \frac{3m^3e}{h^4} (\sin \phi + \phi \cos \phi + \frac{1}{3}e \sin 2\phi).$$

If it were not for the term in  $\phi \cos \phi$ , this would give a perihelion value  $\phi = 2\pi$  as in the Newtonian case. Actually the value is  $\phi = 2\pi + \epsilon$ , where  $\epsilon$  is small. To find  $\epsilon$ , substitute  $2\pi + \epsilon$  for  $\phi$  in the preceding equation. Thus

$$0 = - \frac{me}{h^2} \epsilon + \frac{3m^3e}{h^4} (2\pi),$$

neglecting terms of higher order; so that

$$\epsilon = \frac{6\pi m^2}{h^2}.$$

In the case of the planet Mercury this formula gives for the rotation of the major axis of the orbit in one hundred years approximately 43 sec. of arc.

It has been known to astronomers for a long time that, after allowing for all disturbing influences due to the other planets, there is a rotation of the perihelion of Mercury's orbit of about 43 sec. per

century which cannot be explained by the Newtonian theory. The fact that Einstein's theory gives just this observed rotation is a remarkable success for his theory. For the other planets the correction is inappreciable.

## 12. Deflection of Light Rays by the Sun.

The path of a ray of light in a gravitational field according to Einstein's theory is the same as that of a particle moving with the velocity of light. For such a particle  $ds = 0$  where  $ds$  is any element of its path in the four-dimensional world.

For any particle we have, as already shown,

$$\left(\frac{du}{d\phi}\right)^2 + u^2 = \frac{a^2 - 1}{h^2} + \frac{2mu}{h^2} + 2mu^3,$$

and

$$\frac{d^2u}{d\phi^2} + u = \frac{m}{h^2} + 3mu^2,$$

where  $h = r^2 d\phi/ds$ . For a light ray,  $ds = 0$  and therefore  $h = \infty$ . The differential equation becomes

$$\frac{d^2u}{d\phi^2} + u = 3mu^2.$$

When  $m = 0$ , this is satisfied by

$$u = \frac{1}{p} \cos\phi,$$

the equation of a straight line,  $p$  being the perpendicular from the origin.

Substitute this first approximation as before in the small term on the right. Thus

$$\frac{d^2u}{d\phi^2} + u = \frac{3m}{p^2} \frac{1 + \cos 2\phi}{2},$$

a solution of which is

$$u = \frac{1}{p} \cos\phi + \frac{3m}{p^2} \left(\frac{1}{2} - \frac{1}{6} \cos 2\phi\right),$$

where, as in the first approximation, the angle  $\phi$  is measured from the point where  $u$  is a maximum. This equation will give  $u = 0$  (or  $r = \infty$ ) when  $\phi = \frac{\pi}{2} + \epsilon$  or  $\phi = -\frac{\pi}{2} - \epsilon$ , where  $\epsilon$  is small. To find  $\epsilon$ , we have

$$0 = \frac{1}{p} (-\epsilon) + \frac{3m}{p^2} \left(\frac{1}{2} + \frac{1}{6}\right),$$

or

$$\epsilon = \frac{2m}{p}.$$

The asymptotic directions of the ray therefore deviate by  $\frac{2m}{p}$  radians from the mean direction.

The final deviation of a ray which starts from an infinite distance on one side and travels to an infinite distance on the other side is therefore given by

$$\Delta = \frac{4m}{p}.$$

Thus the light from a star passing by the sun should be deviated at the earth through an angle  $4m/p$ .

We have taken the gravitational potential to be equal to  $m/r$  and the velocity of light to be unity so that our unit of mass is very large.

The gravitational attraction in dynes between two particles of masses  $m$  and  $m'$  gm. at a distance  $r$  cm. apart is  $Gmm'/r^2$  where  $G = 6.66 \times 10^{-8}$ . Hence if  $g$  is the acceleration of a particle at a distance  $r$  from a mass  $m$  then  $g = Gm/r^2$ . If the unit of mass is to be a mass which gives an acceleration equal to the velocity of light in centimetres per second to a particle at a distance from it equal to the number of centimetres light travels in 1 sec. then we have  $3 \times 10^{10} = Gm/9 \times 10^{20}$ , where  $m$  is this new unit of mass expressed in grammes. Hence  $m = 27 \times 10^{30} \times 10^8/6.66$ , which is equal to  $4 \times 10^{33}$  gm.

The mass of the sun is nearly  $2 \times 10^{33}$  gm. or  $\frac{2 \times 10^{33}}{4 \times 10^{33}} = 5 \times 10^{-6}$  in the new unit, and its radius is  $7 \times 10^{10}$  cm. or 2.33 when the unit of length is  $3 \times 10^{10}$  cm. Hence for a ray of light which just grazes the sun's surface the deviation  $\Delta$  should be

$$\frac{4 \times 5 \times 10^{-6}}{2.33} = 8.6 \times 10^{-6}$$

or  $\Delta = 1.77$  sec. of arc.

The apparent positions of several stars when near the sun have been determined by photographing them during a total solar eclipse, and it has been found that there is a deviation of the light approximately equal to that given by Einstein's formula  $\Delta = 4m/p$ .

### 13. Displacement of Spectral Lines.

In the four-dimensional world the interval  $ds$  between two events is supposed to be the same in any co-ordinate system. Also all atoms of the same kind are supposed to be identical in structure, so that the interval  $ds$  between the emission of two successive light waves by an atom should be the same for all such atoms. The presence of a gravitational field should not alter  $ds$  because, as we have seen, the effects due to gravitation are the same as those due to changing co-ordinates.

Consider then two identical atoms, one at the surface of the sun and one on the earth. Let them both be at rest so that  $dx = dy = dz = 0$ . Then if  $ds$  is the interval between the emission of two light waves by either atom we have

$$ds^2 = \left(1 - \frac{2m}{r}\right) dt'^2$$

for the atom on the sun, and  $ds^2 = dt^2$  for the atom on the earth, since  $2m/r$  is negligible for the earth. Hence

$$\frac{dt}{dt'} = \sqrt{1 - \frac{2m}{r}} = 1 - \frac{m}{r}.$$

Thus  $dt'$  is slightly longer than  $dt$ , so that the spectral lines emitted by elements in the sun should be shifted slightly towards the red end of the spectrum.  $m/r$  is equal to  $2.1 \times 10^{-6}$  so that the change of wavelength is very small, only a few thousandths of an Ångström unit. As the result of very careful measurements mainly by C. E. St. John it is now believed that this minute effect actually exists, in agreement with Einstein's theory.

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## CHAPTER XX

### Mathematical Notes

#### 1. Vector Analysis.

Many physical quantities are *vectors*, that is, quantities which have direction as well as magnitude, for example, velocity, acceleration, force, momentum, angular velocity. Quantities which are determined by a number alone without any reference to direction are called *scalars*, for example, mass, energy, electric charge. Two vectors are equal if they have equal numerical magnitudes and are in the same direction. Vectors can be represented by straight lines. The length of the line representing a vector is taken equal to the numerical magnitude of the vector, and the line is drawn parallel to the direction of the vector. It is necessary to regard the line as drawn *from* a point  $A$  to another point  $B$ , and the direction of the vector represented is from  $A$  to  $B$ ; or an arrow head may be put on the line to indicate the direction of the vector along the line, as  $\overrightarrow{AB}$ . The vector  $\mathbf{V}$  (in modern works usually printed thus in heavy type) represented by a line from  $A$  to  $B$  is sometimes said to be of opposite *sense* to the vector  $\mathbf{V}'$  represented by the line from  $B$  to  $A$ . The word "sense", however, seems inappropriate, and we shall say that two such vectors are of opposite *sign*, so that  $\mathbf{V} = -\mathbf{V}'$ .

If  $\mathbf{A}$  and  $\mathbf{B}$  are two vectors, and  $\mathbf{A} = \mathbf{B}$ , then this vector equation means that the magnitudes of  $\mathbf{A}$  and  $\mathbf{B}$  are equal, and also that their directions are the same, so that they could both be represented by the same line drawn from one point to another.

If a vector  $\mathbf{V}$  is represented by a line  $AB$  and another vector  $\mathbf{U}$  by a line  $BC$ , then the vector  $\mathbf{R}$  represented by the line  $AC$  is called the sum or resultant of the vectors  $\mathbf{V}$  and  $\mathbf{U}$ , and this is expressed by the vector equation

$$\mathbf{R} = \mathbf{V} + \mathbf{U}.$$

The difference between  $\mathbf{V}$  and  $\mathbf{U}$  is taken to be the sum of  $\mathbf{V}$  and  $-\mathbf{U}$ , got by drawing a line  $AB$  to represent  $\mathbf{V}$  and a line  $BD$  equal but in the opposite direction to  $BC$  which represents  $\mathbf{U}$ , and taking  $AD$  to represent the difference  $\mathbf{V} - \mathbf{U}$ .

Relative to rectangular axes  $x, y, z$  any vector  $\mathbf{V}$  has three components which may be denoted by  $V_x, V_y, V_z$ . These components are scalar quantities. A vector of magnitude  $V_x$  in the direction of the  $x$  axis may be represented by the symbol  $V_x \mathbf{i}$ , where  $\mathbf{i}$  is a unit vector in the direction  $Ox$ . But it is sometimes convenient for brevity to denote the vector  $V_x \mathbf{i}$  simply by  $\mathbf{V}_x$ . If  $l, m, n$  denote the cosines of the angles between a vector  $\mathbf{V}$  and the axes  $x, y, z$ , then  $V_x = \bar{V}l, V_y = \bar{V}m, V_z = \bar{V}n$ , where  $\bar{V}$  denotes the numerical magnitude of the vector  $\mathbf{V}$ . If the vector  $\mathbf{V}$  is represented by a line  $OP$  drawn from the origin  $O$ , then we can go from  $O$  to  $P$  by moving along the  $x$  axis a distance  $V_x$ , then parallel to the  $y$  axis a distance  $V_y$ , and finally parallel to the  $z$  axis a distance  $V_z$ . Thus we see that  $\bar{V}$  is given by the vector equation

$$\mathbf{V} = \mathbf{V}_x + \mathbf{V}_y + \mathbf{V}_z.$$

If two vectors  $A$  and  $B$  are equal, so that  $\mathbf{A} = \mathbf{B}$ , it follows that  $A_x = B_x, A_y = B_y$ , and  $A_z = B_z$ . Any vector equation is therefore equivalent to three equations between the  $x, y, z$  components of the vectors. For example, if  $\mathbf{R} = \mathbf{V} - \mathbf{U}$  then  $R_x = V_x - U_x, R_y = V_y - U_y, R_z = V_z - U_z$ . The magnitude  $\bar{V}$  of  $\mathbf{V}$  is given by

$$\bar{V}^2 = V_x^2 + V_y^2 + V_z^2.$$

This equation is often written  $V^2 = V_x^2 + V_y^2 + V_z^2$ , because it is clear that only the magnitude of  $\mathbf{V}$  can be involved, so that the bar above the  $V$  may be omitted. The magnitude of a vector  $\mathbf{V}$  is often written  $|\mathbf{V}|$  instead of  $\bar{V}$ . The equation

$$\bar{V}^2 = V_x^2 + V_y^2 + V_z^2$$

with  $V_x = \bar{V}l, V_y = \bar{V}m$ , and  $V_z = \bar{V}n$  gives the well-known result

$$1 = l^2 + m^2 + n^2.$$

If  $l, m, n$  are the direction cosines of  $\mathbf{V}$ , as above, and  $\lambda, \mu, \nu$  the direction cosines of another vector  $\mathbf{U}$ , then the angle  $\theta$  between the two vectors is given by

$$\cos \theta = l\lambda + m\mu + n\nu,$$

for  $\mathbf{V} = \mathbf{V}_x + \mathbf{V}_y + \mathbf{V}_z$ , so that the component of  $\mathbf{V}$  along  $\mathbf{U}$  is equal to the sum of the components of  $\mathbf{V}_x, \mathbf{V}_y$ , and  $\mathbf{V}_z$  along  $\mathbf{U}$ , or  $\lambda V_x + \mu V_y + \nu V_z$ , which is equal to  $l\lambda \bar{V} + m\mu \bar{V} + n\nu \bar{V}$ ; and the component of  $\mathbf{V}$  along  $\mathbf{U}$  is equal to  $\bar{V} \cos \theta$ .

## 2. Scalar Product and Vector Product of Two Vectors.

The product of the magnitude of the component of a vector  $\mathbf{V}$  along another one  $\mathbf{U}$  and the magnitude of  $\mathbf{U}$  is called the *scalar product*, or *inner product*, of  $\mathbf{V}$  and  $\mathbf{U}$ . It may be denoted by  $(\mathbf{V} \cdot \mathbf{U})$ . Thus  $(\mathbf{V} \cdot \mathbf{U}) = \bar{V}\bar{U} \cos \theta$ , where  $\theta$  is the angle between  $\mathbf{U}$  and  $\mathbf{V}$ . Also since  $\bar{V} \cos \theta = \lambda V_x + \mu V_y + \nu V_z$ , and  $U_x = \lambda \bar{U}$ ,  $U_y = \mu \bar{U}$ ,  $U_z = \nu \bar{U}$ , we have

$$(\mathbf{V} \cdot \mathbf{U}) = V_x U_x + V_y U_y + V_z U_z. \quad (1)$$

As an example of a scalar product we may consider the work done by a force  $\mathbf{F}$  when its point of application is displaced a distance  $\mathbf{D}$ . The work is  $(\mathbf{F} \cdot \mathbf{D}) = \bar{F}\bar{D} \cos \theta = F_x D_x + F_y D_y + F_z D_z$ .

A *rotation* about an axis may be regarded as a vector. The direction of this vector is taken to be that in which an ordinary right-handed screw would move along the axis when subjected to the given rotation. An *angular velocity* is regarded as a vector in the same way.

The *outer product*, or *vector product*, of two vectors  $\mathbf{V}$  and  $\mathbf{U}$  is defined to be a vector of magnitude  $\bar{V}\bar{U} \sin \theta$ , where  $\theta$  is the angle between  $\mathbf{V}$  and  $\mathbf{U}$ , and direction that of the rotation from  $\mathbf{V}$  to  $\mathbf{U}$ , through the angle  $\theta$ . It is denoted by  $[\mathbf{V} \cdot \mathbf{U}]$ . Thus  $[\mathbf{V} \cdot \mathbf{U}]$  is a vector equal to the area of the parallelogram formed by  $\mathbf{V}$  and  $\mathbf{U}$ , and its direction is perpendicular to the plane containing  $\mathbf{V}$  and  $\mathbf{U}$ . The angle  $\theta$  is taken to be the positive angle less than  $180^\circ$  between  $\mathbf{V}$  and  $\mathbf{U}$ . The vector  $[\mathbf{V} \cdot \mathbf{U}]$  is of opposite sign to the vector  $[\mathbf{U} \cdot \mathbf{V}]$ , since the direction of rotation is reversed, so that

$$[\mathbf{V} \cdot \mathbf{U}] + [\mathbf{U} \cdot \mathbf{V}] = 0.$$

As an example of a vector product we may take the force per unit length on a current  $\mathbf{C}$  in a magnetic field  $\mathbf{H}$ , which is equal to  $CH \sin \theta$ , and is perpendicular to  $\mathbf{C}$  and  $\mathbf{H}$ . It is equal to  $[\mathbf{C} \cdot \mathbf{H}]$ .

The components of the vector product are equal to the projections of the area of the parallelogram formed by  $\mathbf{V}$  and  $\mathbf{U}$  on the planes perpendicular to the axes. This projection on the  $xy$  plane is equal to the area of the parallelogram formed by the points  $(0, 0)$ ,  $(U_x, U_y)$ , and  $(V_x, V_y)$ , which is easily seen to be equal to  $V_x U_y - V_y U_x$ . The components of the vector product are therefore given by

$$\left. \begin{aligned} [\mathbf{V} \cdot \mathbf{U}]_x &= V_y U_z - V_z U_y, \\ [\mathbf{V} \cdot \mathbf{U}]_y &= V_z U_x - V_x U_z, \\ [\mathbf{V} \cdot \mathbf{U}]_z &= V_x U_y - V_y U_x. \end{aligned} \right\} \quad (2)$$

Another example of a vector product is the moment of a force  $\mathbf{F}$  acting at a point  $P$  with co-ordinates  $x, y, z$  about an axis through the origin

perpendicular to the plane containing the force and the origin. This is equal to  $\bar{F}r \sin\theta$ , where  $r = OP$ , and  $\theta$  is the angle between  $OP$  produced and  $\mathbf{F}$ . The  $x$  component of this vector product is the moment of  $\mathbf{F}$  about the  $x$  axis, which is equal to  $yF_z - zF_y$ , so that

$$yF_z - zF_y = [\mathbf{r} \cdot \mathbf{F}]_x.$$

If  $\mathbf{U}$ ,  $\mathbf{V}$ ,  $\mathbf{W}$  are three vectors then  $(\mathbf{U} \cdot [\mathbf{V} \cdot \mathbf{W}])$  can easily be seen to be the volume of the parallelepiped formed by the three vectors. Hence if  $(\mathbf{U} \cdot [\mathbf{V} \cdot \mathbf{W}]) = 0$  the three vectors must lie in the same plane.

### 3. Scalar and Vector Fields.

The *field* of a vector or scalar quantity is a region throughout which at every point the quantity has a definite value, which in general varies from point to point. For example, the space around electrically charged bodies is a *vector field* of the electric field strength, and the atmosphere is a *scalar field* of the density of the air.

In the field of a scalar  $S$  the vector called the *gradient* of  $S$  is defined to be a vector the components of which are  $\frac{\partial S}{\partial x}$ ,  $\frac{\partial S}{\partial y}$ , and  $\frac{\partial S}{\partial z}$ ; or, if at any point in the field we draw a surface on which  $S$  is constant, then the gradient of  $S$  is normal to the surface, and its magnitude is the rate of variation of  $S$  per unit length along the normal. If  $p$  denotes the distance measured along a curve which is normal to the surfaces of constant  $S$ , then  $\text{grad } S = \frac{\partial S}{\partial p}$ , and  $(\text{grad } S)_x = \frac{\partial S}{\partial x}$ ,  $(\text{grad } S)_y = \frac{\partial S}{\partial y}$ ,  $(\text{grad } S)_z = \frac{\partial S}{\partial z}$ , so that we have the vector equation,  $\text{grad } S = \frac{\partial S}{\partial p}$ , and also

$$|\text{grad } S|^2 = \left(\frac{\partial S}{\partial p}\right)^2 = \left(\frac{\partial S}{\partial x}\right)^2 + \left(\frac{\partial S}{\partial y}\right)^2 + \left(\frac{\partial S}{\partial z}\right)^2.$$

It is evident that the component of  $\text{grad } S$  in any direction is equal to  $\frac{\partial S}{\partial s}$ , where  $\partial s$  is an infinitesimal displacement in the given direction.

### 4. Line Integrals. Potentials.

Consider a curve of any shape drawn in a scalar or vector field. Let the distance measured along this curve be  $s$ , so that  $ds$  is an element of the curve. If  $Q$  denotes any quantity which has a definite value at every point in the field, then the sum of the products of the lengths of the elements  $ds$  of the curve and the values of  $Q$  at these elements, or  $\int Q ds$ , is called the line integral of  $Q$  along the curve. The line integral

for the part of the curve between two points  $A$  and  $B$  on it is denoted by  $\int_A^B Q ds$ . If  $\mathbf{V}$  is a vector, and  $V_s$  denotes its component along the curve, then  $\int_A^B V_s ds$  is the line integral of  $V_s$  along the curve. The product  $V_s ds$  is equal to the scalar product of  $\mathbf{V}$  and  $d\mathbf{s}$ , so that

$$\int_A^B V_s ds = \int_A^B (\mathbf{V} \cdot d\mathbf{s}) = \int_A^B (V_x dx + V_y dy + V_z dz),$$

where  $dx, dy, dz$  are the components of  $d\mathbf{s}$ . For the gradient of a scalar  $S$ , or  $\text{grad } S$ , we have therefore

$$\int_A^B (\text{grad } S)_s ds = \int_A^B \frac{\partial S}{\partial s} ds,$$

so that

$$\int_A^B (\text{grad } S)_s ds = S_B - S_A.$$

Thus the value of the integral  $\int_A^B (\text{grad } S)_s ds$  between two points  $A$  and  $B$  depends only on the values of  $S$  at  $A$  and at  $B$ , and is independent of the position of the curve joining  $A$  and  $B$  along which the integral is taken.

If a vector  $\mathbf{V}$  is equal to  $(-\text{grad } S)$ , where  $S$  is a scalar, then  $S$  is called the potential of  $\mathbf{V}$ . Hence

$$\int_A^B V_s ds = - \int_A^B (\text{grad } S)_s ds = S_A - S_B,$$

that is, the potential difference between two points in the vector field is equal to the line integral of the vector component along any curve joining the points. The components of the vector are equal to the components of  $(-\text{grad } S)$ , so that

$$V_x = -\frac{\partial S}{\partial x}, \quad V_y = -\frac{\partial S}{\partial y}, \quad V_z = -\frac{\partial S}{\partial z}.$$

Thus if we know the potential of a vector at all points in the field we can easily calculate the vector at any point.

Examples of potentials are electrostatic potential, magnetostatic potential, and gravitational potential.

### 5. Vector Lines and Tubes.

A line drawn in the field of a vector  $\mathbf{V}$  so as to be everywhere along the direction of  $\mathbf{V}$  is called a *vector line*.

If we take a very small closed curve in the field and draw vector lines through all the points on it these lines enclose a tube called a *vector tube*.

The magnitude of the vector at a particular cross-section of a vector tube can be represented by drawing vector lines inside the tube

uniformly distributed over the cross-section, so that the number of the lines is proportional to the product of the magnitude of the vector and the area of the cross-section. Thus if  $\alpha$  denotes the area of a cross-section of the tube and  $\bar{V}$  the magnitude of the vector at this cross-section, then we suppose the number of vector lines drawn in the tube to be proportional to  $\alpha\bar{V}$ . The number of lines per unit area of cross-section is then proportional to  $\bar{V}$ . In a very important class of cases the number of lines is constant from one cross-section to another, but this is not so in general. See §6, 13 below.

Instead of supposing the magnitude of the vector represented in this way by the number of vector lines drawn in the field, we may suppose the vector tube divided into a number of equal smaller tubes proportional to the product  $\alpha V$ , and so have the number of tubes per unit area proportional to the magnitude of the vector. In this case  $V$  is inversely proportional to the cross-section of the vector tubes.

If the tubes are drawn so that the number per unit area on a surface perpendicular to the vector  $\mathbf{V}$  is equal to the magnitude of  $\mathbf{V}$ , the tubes are called unit tubes of the vector  $\mathbf{V}$ . The cross-section of a unit tube at any point is then equal to  $1/\bar{V}$ . The field is not well represented in this way unless the tubes are of small cross-section, but, of course, we may if necessary divide the unit tubes into as many smaller tubes as we like each of cross-section  $1/n\bar{V}$ , where  $n$  is any convenient number.

The unit tubes may be closed tubes, or they may start at one point and end at another in the field. The number of unit tubes which start per unit volume at a point in the field minus the number which end per unit volume at the point is called the divergence of the vector  $\mathbf{V}$  and is written  $\text{div } \mathbf{V}$ . The divergence of  $\mathbf{V}$  may also be defined by the equation

$$\text{div } \mathbf{V} = \frac{\partial V_x}{\partial x} + \frac{\partial V_y}{\partial y} + \frac{\partial V_z}{\partial z} \quad \dots \quad (3)$$

That the two definitions are equivalent will be shown in the next section.

## 6. Green's Theorem.

An important theorem giving transformations of volume integrals into surface integrals was discovered by Green. A special case discovered earlier by Gauss is known as Gauss's theorem.

Consider a closed surface of area  $\sigma$  enclosing a volume  $S$ . Let  $A$  be any function of position which has a definite value at all points inside the closed surface, and consider the integral  $\int \frac{\partial A}{\partial x} dx dy dz$  taken

over the volume  $S$ . Draw a line through the volume  $S$  parallel to the  $x$  axis and let it cut the closed surface at two points  $P$  and  $Q$ . The line integral  $\int \frac{\partial A}{\partial x} dx$  along this line from  $P$  to  $Q$  is equal to  $A_Q - A_P$ .

If we multiply this by the small rectangular area  $dy dz$  we get therefore the volume integral of  $\partial A / \partial x$  over the volume of a rod of uniform cross-section  $dy dz$  and length  $x_Q - x_P$ . Let  $d\sigma$  be the area cut off on the surface  $\sigma$  by the rod at  $Q$ . Then if  $l$  denotes the cosine of the angle between the outward drawn normal to  $\sigma$  at  $Q$  and the  $x$  axis we have  $dy dz = l d\sigma$ , since  $dy dz$  is equal to the projection of  $d\sigma$  on the  $yz$  plane. Hence  $A_Q dy dz = l A_Q d\sigma$ . In the same way if  $d\sigma'$  is the area cut off at  $P$  and  $l'$  the cosine of the angle between the outward normal to  $\sigma$  at  $P$  and the  $x$  axis, then

$$A_P dy dz = -l' A_P d\sigma'.$$

Hence

$$(A_Q - A_P) dy dz = l A_Q d\sigma + l' A_P d\sigma'.$$

Now the whole volume  $S$  may be divided up into small elements like  $(x_Q - x_P) dy dz$ , and for each of these the volume integral of  $\partial A / \partial x$  is equal to an expression like that just found. We see therefore that the volume integral of  $\partial A / \partial x$  for the whole volume  $S$  is equal to the sum of all these quantities like  $l A_Q d\sigma + l' A_P d\sigma'$ . But this sum is equal to the integral of  $l A d\sigma$  over the whole surface  $\sigma$ , since when the whole volume  $S$  is divided into rod elements the ends of the elements will cover the whole surface  $\sigma$ . Hence we get

$$\int_S \frac{\partial A}{\partial x} dx dy dz = \int_{\sigma} l A d\sigma.$$

In the same way, if  $m$  and  $n$  are the cosines of the angle between an outward drawn normal to  $\sigma$  and the  $y$  and  $z$  axes, and  $B$  and  $C$  any functions of position inside the volume  $S$ , then

$$\int_S \frac{\partial B}{\partial y} dx dy dz = \int_{\sigma} m B d\sigma,$$

and

$$\int_S \frac{\partial C}{\partial z} dx dy dz = \int_{\sigma} n C d\sigma.$$

Adding the three results we get

$$\int_S \left( \frac{\partial A}{\partial x} + \frac{\partial B}{\partial y} + \frac{\partial C}{\partial z} \right) dx dy dz = \int_{\sigma} (l A + m B + n C) d\sigma, \quad (4)$$

which is Green's theorem.

If

$$A = V_x, \quad B = V_y, \quad C = V_z,$$

where  $\mathbf{V}$  is a vector, then

$$lV_x + mV_y + nV_z = V_n,$$

where  $V_n$  denotes the component of  $\mathbf{V}$  along the outward normal to  $d\sigma$ , so that

$$\int_s \left( \frac{\partial V_x}{\partial x} + \frac{\partial V_y}{\partial y} + \frac{\partial V_z}{\partial z} \right) dx dy dz = \int_\sigma V_n d\sigma. \quad . \quad . \quad (5)$$

Now since the normal cross-section of a unit vector tube is  $1/\bar{V}$ , the section by a surface the normal to which makes an angle  $\theta$  with  $V$  is  $1/V \cos \theta$ . Thus, if  $V_n = V \cos \theta$ , we see that the number of unit tubes leaving the closed surface through the element of area  $d\sigma$  is  $V \cos \theta d\sigma = V_n d\sigma$ . Hence  $\int V_n d\sigma$  is equal to the total number of unit vector tubes coming out of the volume  $S$  through the surface  $\sigma$ . Tubes entering the surface are here understood to be reckoned negative.

Thus it appears that

$$\int_s \left( \frac{\partial V_x}{\partial x} + \frac{\partial V_y}{\partial y} + \frac{\partial V_z}{\partial z} \right) dx dy dz$$

is equal to the total number of unit vector tubes leaving the volume  $S$ . This is true for any volume, so that if we consider a very small one we see that

$$\frac{\partial V_x}{\partial x} + \frac{\partial V_y}{\partial y} + \frac{\partial V_z}{\partial z}$$

is equal to the number of unit tubes which start from it per unit volume. We have therefore

$$\text{div } \mathbf{V} = \frac{\partial V_x}{\partial x} + \frac{\partial V_y}{\partial y} + \frac{\partial V_z}{\partial z}, \quad . \quad . \quad . \quad (3')$$

and so, putting  $dx dy dz = dS$ , we get

$$\int_s \text{div } \mathbf{V} dS = \int_\sigma V_n d\sigma. \quad . \quad . \quad . \quad (6)$$

This result is often called Gauss's theorem.

Green's theorem enables a volume integral of any function of  $x$ ,  $y$ , and  $z$  which can be expressed in the form

$$\frac{\partial A}{\partial x} + \frac{\partial B}{\partial y} + \frac{\partial C}{\partial z}$$

to be transformed into a surface integral over the surface enclosing the volume.

The volume  $S$  may be the space between two closed surfaces, one inside the other, in which case the surface integral is taken over both

surfaces, and the normals are drawn in both cases in the direction from the enclosed volume to the surface.

### 7. Another Form of Green's Theorem.

$$\text{Let} \quad A = \phi \frac{\partial \psi}{\partial x}, \quad B = \phi \frac{\partial \psi}{\partial y}, \quad C = \phi \frac{\partial \psi}{\partial z},$$

so that

$$\frac{\partial A}{\partial x} + \frac{\partial B}{\partial y} + \frac{\partial C}{\partial z} = \phi \left( \frac{\partial^2 \psi}{\partial x^2} + \frac{\partial^2 \psi}{\partial y^2} + \frac{\partial^2 \psi}{\partial z^2} \right) + \frac{\partial \phi}{\partial x} \frac{\partial \psi}{\partial x} + \frac{\partial \phi}{\partial y} \frac{\partial \psi}{\partial y} + \frac{\partial \phi}{\partial z} \frac{\partial \psi}{\partial z}.$$

If  $\frac{\partial^2 \psi}{\partial x^2} + \frac{\partial^2 \psi}{\partial y^2} + \frac{\partial^2 \psi}{\partial z^2}$  be denoted by  $\Delta \psi$ , Green's theorem now gives

$$\begin{aligned} \int_S (\phi \Delta \psi + (\text{grad } \phi \cdot \text{grad } \psi)) dS \\ = \int_\sigma \phi \left( l \frac{\partial \psi}{\partial x} + m \frac{\partial \psi}{\partial y} + n \frac{\partial \psi}{\partial z} \right) d\sigma = \int_\sigma \phi \frac{\partial \psi}{\partial n} d\sigma. \end{aligned}$$

In the same way, interchanging  $\phi$  and  $\psi$ , we get

$$\int_S (\psi \Delta \phi + (\text{grad } \phi \cdot \text{grad } \psi)) dS = \int_\sigma \psi \frac{\partial \phi}{\partial n} d\sigma.$$

Thus, by subtraction,

$$\int_S (\phi \Delta \psi - \psi \Delta \phi) dS = \int_\sigma \left( \phi \frac{\partial \psi}{\partial n} - \psi \frac{\partial \phi}{\partial n} \right) d\sigma. \quad (7)$$

### 8. Solution of Poisson's Equation.

As an example of the use of this form of Green's theorem, we may find the solution of the equation

$$\frac{\partial^2 A}{\partial x^2} + \frac{\partial^2 A}{\partial y^2} + \frac{\partial^2 A}{\partial z^2} = \omega, \quad (8)$$

or  $\Delta A = \omega$ , where  $A$  and  $\omega$  are scalar functions of  $x$ ,  $y$ , and  $z$ .

Put  $\psi = A$  and  $\phi = 1/r$ , where  $r^2 = x^2 + y^2 + z^2$ , so that

$$\Delta \phi = -\frac{3}{r^3} + \frac{3(x^2 + y^2 + z^2)}{r^5} = 0. \quad \checkmark$$

Hence

$$\int_S \frac{\omega}{r} dS = \int_\sigma \left( \frac{1}{r} \frac{\partial A}{\partial n} + \frac{A}{r^2} \frac{\partial r}{\partial n} \right) d\sigma.$$

Now let  $S$  be the volume between two concentric spheres of radii  $r_1$  and  $r_2$ , with centre at the origin. In this case  $\frac{\partial A}{\partial n} = \frac{\partial A}{\partial r}$  on the outer

sphere  $r_2$ , and  $\frac{\partial A}{\partial n} = -\frac{\partial A}{\partial r}$  on the inner one  $r_1$ . Also  $\frac{\partial r}{\partial n} = 1$  on the outer, and  $-1$  on the inner, sphere. Therefore

$$\int_s \frac{\omega}{r} dS = \int_{\sigma_2} \left( \frac{1}{r_2} \frac{\partial A}{\partial r} + \frac{A}{r_2^2} \right) d\sigma - \int_{\sigma_1} \left( \frac{1}{r_1} \frac{\partial A}{\partial r} + \frac{A}{r_1^2} \right) d\sigma.$$

Now let the outer sphere be very large, and assume that  $A$  and  $\partial A/\partial r$  become very small when  $r$  becomes very great, in such a way that the surface integral over the outer sphere tends to zero when  $r_2$  is indefinitely increased. Also let the inner sphere be very small, so that  $A$  may be considered to be constant over its surface. Then

$$\int_s \frac{\omega}{r} dS = -4\pi r_1^2 \left( \frac{1}{r_1} \frac{\partial \bar{A}}{\partial r} + \frac{\bar{A}}{r_1^2} \right),$$

where  $\frac{\partial \bar{A}}{\partial r}$  denotes the average value of  $\partial A/\partial r$  over the small sphere.

Taking the limit for  $r_1 = 0$ , we have

$$\int_s \frac{\omega}{r} dS = -4\pi A,$$

or

$$A = -\frac{1}{4\pi} \int_s \frac{\omega}{r} dS. \quad . \quad . \quad . \quad . \quad . \quad (9)$$

Here  $-\frac{1}{4\pi} \int_s \frac{\omega}{r} dS$  is the value of  $A$  at the origin  $r = 0$ . The origin, of course, can be taken to be anywhere, so that we see that if

$$\Delta A = \omega,$$

then the value of  $A$  at any point can be obtained by dividing the space around the point into small elements of volume  $dS$  and multiplying each element by the value of  $\omega/r$  at the element, where  $r$  is the distance of the element  $dS$  from the point at which  $A$  is required. The sum of all these products multiplied by  $-1/4\pi$ , i.e.  $-\frac{1}{4\pi} \int \frac{\omega dS}{r}$ , is then equal to  $A$ .

If, for example,  $\omega = 0$  everywhere except in a small volume  $v$  in which its average value is  $\bar{\omega}$ , and if  $v\bar{\omega} = e$ , then at any point at a distance  $r$  from this small volume we have  $A = -e/4\pi r$ .

In the equations  $\Delta A = \omega$  and  $A = -\frac{1}{4\pi} \int_s \frac{\omega}{r} dS$ ,  $A$  and  $\omega$  may be the components of vector quantities. Thus if  $\mathbf{A}$  and  $\boldsymbol{\omega}$  are vectors, and  $\Delta \mathbf{A} = \boldsymbol{\omega}$ , then

$$A_x = -\frac{1}{4\pi} \int \frac{\omega_x}{r} dS,$$

with similar equations for  $A_y$  and  $A_z$ .

Thus we see that  $\Delta \mathbf{A} = \boldsymbol{\omega}$ , where  $\mathbf{A}$  and  $\boldsymbol{\omega}$  are vectors, gives

$$\mathbf{A} = -\frac{1}{4\pi} \int \frac{\boldsymbol{\omega} dS}{r}$$

where  $\int \frac{\boldsymbol{\omega} dS}{r}$  now denotes the vector sum of the products  $\boldsymbol{\omega} dS/r$ . We may regard the vector field of  $\mathbf{A}$  as excited or produced by the quantity  $\boldsymbol{\omega}$ .

### 9. Solution of Equation for a Propagated Potential.

Consider next the equation

$$\Delta A - \frac{1}{c^2} \ddot{A} = \omega, \quad . . . . . (10)$$

where  $c$  is a constant and  $\ddot{A} = \partial^2 A / \partial t^2$ . This important equation can be regarded as meaning that  $\omega$  excites a field which is propagated out from  $\omega$  with the velocity  $c$ .

The solution of this equation can be obtained by means of Green's theorem in a manner similar to that used to get the solution of  $\Delta A = \omega$ .

Put  $\phi = \frac{1}{r} F(t + r/c)$ , and  $\psi = A$ , in equation (7):

$$\int_s (\phi \Delta \psi - \psi \Delta \phi) dS = \int_\sigma \left( \phi \frac{\partial \psi}{\partial n} - \psi \frac{\partial \phi}{\partial n} \right) d\sigma.$$

Here  $F(t + r/c)$  denotes an arbitrary function of  $t + r/c$ , which can be given any desired form. It can easily be shown that  $\phi = \frac{1}{r} F(t + r/c)$  satisfies  $\Delta \phi - \frac{1}{c^2} \ddot{\phi} = 0$ , by substituting the values of the differential coefficients of  $\phi$  in this equation. We have therefore

$$\int_s \left( \phi \left( \omega + \frac{1}{c^2} \ddot{A} \right) - A \frac{1}{c^2} \ddot{\phi} \right) dS = \int_\sigma \left( \phi \frac{\partial \psi}{\partial n} - \psi \frac{\partial \phi}{\partial n} \right) d\sigma,$$

$$\text{or} \quad \int_s \phi \omega dS + \frac{d}{dt} \int_s \frac{(\phi \dot{A} - A \dot{\phi}) dS}{c^2} = \int_\sigma \left( \phi \frac{\partial \psi}{\partial n} - \psi \frac{\partial \phi}{\partial n} \right) d\sigma.$$

Multiply by  $dt$  and integrate from  $t_1$  to  $t_2$ . Thus

$$\int_{t_1}^{t_2} dt \int_s \phi \omega dS + \left[ \int_s \frac{(\phi \dot{A} - A \dot{\phi}) dS}{c^2} \right]_{t_1}^{t_2} = \int_{t_1}^{t_2} dt \int_\sigma \left( \phi \frac{\partial \psi}{\partial n} - \psi \frac{\partial \phi}{\partial n} \right) d\sigma. \quad (11)$$

Now suppose that  $F(t + r/c) = 0$  except for values of  $t + r/c$  very nearly  $= 0$ . Also let  $t_2$  be greater than  $-r/c$ , and  $t_1$  less, so that

the instant  $t = -r/c$  is between  $t_1$  and  $t_2$ . Also suppose that  $\int_{t_1}^{t_2} F(t+r/c) dt = 1$ . This requires that  $F(0)$  be so large that although  $F(t+r/c)$  is zero for all values of  $t+r/c$ , except those near zero, yet  $\int_{t_1}^{t_2} F(t+r/c) dt = 1$ . The function  $F(t+r/c)$  is thus made to act as a selecting factor, since when multiplied by any quantity and integrated over the time it selects the value of the quantity at the time  $t = -r/c$ .

Thus

$$\int_{t_1}^{t_2} dt \int_S \phi \omega dS = \int \frac{[\omega]}{r} dS,$$

where  $[\omega]$  denotes the value of  $\omega$  in the element of volume  $dS$  at the time  $t = -r/c$ .

Hence

$$\int_S \frac{[\omega]}{r} dS = \int_{t_1}^{t_2} dt \int_\sigma \left( \phi \frac{\partial A}{\partial n} - A \frac{\partial \phi}{\partial n} \right) d\sigma,$$

the second term on the left of equation (11) disappearing, since  $\phi$  and  $\dot{\phi}$  vanish when  $t = t_1$  and  $t = t_2$ . Now let the boundary of  $S$  be a very small sphere of radius  $r_1$  and a large sphere of radius  $r_2$ , both with centre at the origin. Assume that  $A$  and  $\partial A/\partial n$  are zero on the large sphere, so that only the integral over the surface of the small sphere need be considered.

We have  $\frac{\partial A}{\partial n} = -\frac{\partial A}{\partial r}$  and  $\frac{\partial \phi}{\partial n} = -\frac{\partial \phi}{\partial r}$ ; also

$$\frac{\partial \phi}{\partial r} = \frac{\partial}{\partial r} \left( \frac{1}{r} F(t+r/c) \right) = -\frac{1}{r_1^2} F + \frac{1}{r_1 c} F',$$

where  $F'$  is the differential coefficient of  $F$  with respect to  $t+r/c$ . Hence

$$\phi \frac{\partial A}{\partial n} - A \frac{\partial \phi}{\partial n} = -\frac{F}{r_1} \frac{\partial A}{\partial r} - \frac{AF}{r_1^2} + \frac{AF'}{r_1 c}.$$

When  $r_1$  is very small, we may regard  $A$  and  $F$  as constant over the surface of the small sphere, and replace  $\partial A/\partial r$  and  $F'$  by their average values, so that

$$\int_{t_1}^{t_2} dt \int_\sigma \left( \phi \frac{\partial A}{\partial n} - A \frac{\partial \phi}{\partial n} \right) d\sigma = \int_{t_1}^{t_2} 4\pi r_1^2 \left( \frac{\overline{AF'}}{r_1 c} - \frac{AF}{r_1^2} - \frac{F}{r_1} \frac{\partial A}{\partial r} \right) dt.$$

In the limit when  $r_1 = 0$ , this becomes

$$-4\pi \int_{t_1}^{t_2} AF dt,$$

which is equal to the value of  $-4\pi A$  at the time  $t = 0$ , because  $F$  is zero except at  $t = -r/c$ , i.e. when  $r = 0$ , at  $t = 0$ .

Hence we have finally

$$A_{t=0} = -\frac{1}{4\pi} \int_s \frac{[\omega] dS}{r}, \quad . . . . . (12)$$

where  $[\omega]$  denotes the value of  $\omega$  in the element of volume  $dS$  at the time  $t = -r/c$ , and  $r$  is the distance from  $dS$  to the point at which  $A$  is to be calculated.

Thus we see that the field at any time due to the  $\omega$  in any element of volume is determined by the value of  $\omega$  at a time  $r/c$  previously, so that the field excited by  $\omega$  travels out with the velocity  $c$ .

### 10. Curl of a Vector.

In the field of any vector  $\mathbf{V}$  we can define another vector  $\mathbf{R}$  determined by the variation of  $\mathbf{V}$  in the following way. Consider a small plane area  $\alpha$  enclosed by a curve of length  $s$ . Take the line integral  $\int V_s ds$  of the component  $V_s$  of  $\mathbf{V}$  along  $s$  round the curve  $s$ . Then  $R_n = \frac{1}{\alpha} \int_s \mathbf{V}_s ds$  can be proved to be the component of a vector  $\mathbf{R}$  along the normal  $n$  to the plane area  $\alpha$ . The direction of  $\mathbf{R}$  along the normal to  $\alpha$  is taken so that a right-handed screw would advance in the direction of  $\mathbf{R}$  when turned in the direction in which the line integral round  $s$  is taken. The vector  $\mathbf{R}$  is called the *curl* (or *rotation*) of  $\mathbf{V}$  and is denoted by  $\text{curl } \mathbf{V}$  (or  $\text{rot } \mathbf{V}$ ).

### 11. Stokes's Theorem.

We shall now consider an important theorem due to Sir G. G. Stokes, which enables an integral over a surface to be transformed into a line integral round the boundary of the surface.

Consider a surface of any shape in the field of a vector  $\mathbf{V}$  and suppose a closed curve drawn on it. Let  $s$  denote the length of the closed curve and  $\sigma$  the area on the surface enclosed by the curve.

Let  $d\sigma$  be an element of area on the surface,  $n$  the normal to the surface at  $d\sigma$ , and  $ds$  an element of the closed curve. Then Stokes's theorem is:

$$\int_\sigma (\text{curl } \mathbf{V})_n d\sigma = \int_s V_s ds, \quad . . . . . (13)$$

or, in words, the integral of the normal component of the curl of a vector over the surface, is equal to the line integral round the boundary of the component of the vector along the bounding curve.

Let  $ABC$  be the closed curve, and let the surface bounded by it be divided up into a large number of small elements by two sets of curves, as shown in fig. 1.

Consider the sum of the line integrals of  $V_s$  round the boundaries

of all these elements going round them all the same way. Each line element not on the outside boundary will appear twice in this sum, once for each of the two elements of area between which it lies. The two line integrals along any line element not on the boundary are equal but of opposite sign, since the element is traversed in opposite directions when going round the two elements of area on either side of the line. The sum of the line integrals of  $V_s$  round all the elements of area into which the whole area is divided is therefore equal to the line integral of  $V_s$  round the outside boundary, or  $\int_s V_s ds$ .

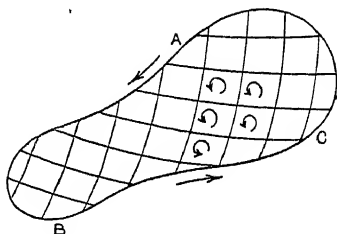


Fig. 1

Now, for any element of area  $d\sigma$ , the line integral of  $V_s$  round its boundary is equal to the component of  $\text{curl } \mathbf{V}$  along the normal to the element multiplied by  $d\sigma$ , or  $(\text{curl } \mathbf{V})_n d\sigma = \int_s V_s ds$ . Hence the sum of all the line integrals round the elements of area is equal to

$$\int_{\sigma} (\text{curl } \mathbf{V})_n d\sigma,$$

so that we get

$$\int_{\sigma} (\text{curl } \mathbf{V})_n d\sigma = \int_s V_s ds,$$

which is Stokes's theorem.

## 12. Components of the Curl of a Vector.

The components of the vector  $\text{curl } \mathbf{V}$  relative to axes  $x, y, z$  can be easily calculated. Consider a small rectangular area parallel to the  $yz$  plane with sides  $dy$  and  $dz$ . The normal to it is along the  $x$  direction, so that we have

$$(\text{curl } \mathbf{V})_x = \frac{1}{dy dz} \int V_s ds,$$

the line integral of  $V_s$  being taken round the rectangular area. Hence

$$\int V_s ds = (V_y - V_y') dy + (V_z' - V_z) dz,$$

where  $V_y, V_z$  and  $V_y', V_z'$  are the values on opposite sides of the area  $dy dz$ .

But  $V_y' - V_y = \frac{\partial V_y}{\partial z} dz$ , and  $V_z' - V_z = \frac{\partial V_z}{\partial y} dy$ , so that

$$\int V_s ds = \left( \frac{\partial V_z}{\partial y} - \frac{\partial V_y}{\partial z} \right) dy dz,$$

and therefore

$$\left. \begin{aligned} (\text{curl } \mathbf{V})_x &= \frac{\partial V_z}{\partial y} - \frac{\partial V_y}{\partial z}, \\ (\text{curl } \mathbf{V})_y &= \frac{\partial V_x}{\partial z} - \frac{\partial V_z}{\partial x}, \\ (\text{curl } \mathbf{V})_z &= \frac{\partial V_y}{\partial x} - \frac{\partial V_x}{\partial y}. \end{aligned} \right\} \quad . \quad . \quad . \quad (14)$$

The curl of a vector may be defined by these equations instead of (§10) by

$$(\text{curl } \mathbf{V})_n = \frac{1}{\alpha} \int V_s ds. \quad . \quad . \quad . \quad (15)$$

### 13. Some Important Relations.

We have

$$\begin{aligned} \text{div} (\text{curl } \mathbf{V}) &= \frac{\partial}{\partial x} \left( \frac{\partial V_z}{\partial y} - \frac{\partial V_y}{\partial z} \right) + \frac{\partial}{\partial y} \left( \frac{\partial V_x}{\partial z} - \frac{\partial V_z}{\partial x} \right) + \frac{\partial}{\partial z} \left( \frac{\partial V_y}{\partial x} - \frac{\partial V_x}{\partial y} \right) \\ &= \frac{\partial^2 V_x}{\partial y \partial z} - \frac{\partial^2 V_x}{\partial z \partial y} + \frac{\partial^2 V_y}{\partial z \partial x} - \frac{\partial^2 V_y}{\partial x \partial z} + \frac{\partial^2 V_z}{\partial x \partial y} - \frac{\partial^2 V_z}{\partial y \partial x}, \end{aligned}$$

$$\text{or} \quad \text{div} (\text{curl } \mathbf{V}) = 0. \quad . \quad . \quad . \quad (16)$$

This means that the vector lines of  $\text{curl } \mathbf{V}$  always form closed curves, so that the number of unit tubes of  $\text{curl } \mathbf{V}$  crossing any section of a vector tube of  $\text{curl } \mathbf{V}$  is constant along the tube. Hence if  $\alpha$  is the area of the cross-section of a vector tube of  $\text{curl } \mathbf{V}$ ,  $\alpha \text{ curl } \mathbf{V}$  is constant along the tube. This is true for any vector the divergence of which is zero; it follows at once from Green's theorem (5).

If the curl of a vector  $\mathbf{V}$  is zero throughout a region in the field of the vector, the line integral of  $V_s$  round any closed curve  $s$  in this region is zero. The vector therefore has a potential, or is equal to the negative gradient of a scalar. The converse is proved as follows:

$$\begin{aligned} (\text{curl} (\text{grad } S))_x &= \frac{\partial}{\partial y} (\text{grad } S)_z - \frac{\partial}{\partial z} (\text{grad } S)_y \\ &= \frac{\partial^2 S}{\partial y \partial z} - \frac{\partial^2 S}{\partial z \partial y} = 0, \end{aligned}$$

with similar equations for the  $y$  and  $z$  components, so that

$$\text{curl} (\text{grad } S) = 0. \quad . \quad . \quad . \quad (17)$$

The field of a vector the curl of which is zero is said to be *irrotational*, and that of a vector with zero divergence is called *solenoidal*.

The curl of the curl of a vector is an important quantity. We have

$$\begin{aligned}
 (\text{curl } (\text{curl } \mathbf{V}))_x &= \frac{\partial}{\partial y} (\text{curl } \mathbf{V})_z - \frac{\partial}{\partial z} (\text{curl } \mathbf{V})_y \\
 &= \frac{\partial^2 V_y}{\partial y \partial x} - \frac{\partial^2 V_x}{\partial y^2} - \frac{\partial^2 V_x}{\partial z^2} + \frac{\partial^2 V_z}{\partial z \partial x} \\
 &= \frac{\partial}{\partial x} \left( \frac{\partial V_x}{\partial x} + \frac{\partial V_y}{\partial y} + \frac{\partial V_z}{\partial z} \right) - \left( \frac{\partial^2 V_x}{\partial x^2} + \frac{\partial^2 V_x}{\partial y^2} + \frac{\partial^2 V_x}{\partial z^2} \right) \\
 &= (\text{grad div } \mathbf{V})_x - \Delta V_x,
 \end{aligned}$$

with similar equations for the  $y$  and  $z$  components. Hence

$$\text{curl curl } \mathbf{V} = \text{grad div } \mathbf{V} - \Delta \mathbf{V}. \quad . \quad . \quad . \quad (18)$$

We have seen (§8) that if  $\Delta \mathbf{V} = \boldsymbol{\omega}$ , then

$$\mathbf{V} = -\frac{1}{4\pi} \int \frac{\boldsymbol{\omega} dS}{r},$$

so that, since

$$\Delta \mathbf{V} = \text{grad div } \mathbf{V} - \text{curl curl } \mathbf{V}$$

we have, for any vector  $\mathbf{V}$ ,

$$\mathbf{V} = \frac{1}{4\pi} \int \frac{1}{r} \text{curl curl } \mathbf{V} dS - \frac{1}{4\pi} \int \frac{1}{r} \text{grad div } \mathbf{V} dS. \quad . \quad (19)$$

If the divergence of a vector  $\mathbf{V}$  is equal to zero, it is always possible to find another vector  $\mathbf{A}$  such that  $\mathbf{V} = \text{curl } \mathbf{A}$ . It can be shown in fact that if  $\mathbf{A} = \frac{1}{4\pi} \int \frac{1}{r} \text{curl } \mathbf{V} dS$ , then  $\mathbf{V} = \text{curl } \mathbf{A}$ .

In the same way, if  $\text{curl } \mathbf{V} = 0$ , it is always possible to find a scalar  $\phi$  such that  $\mathbf{V} = -\text{grad } \phi$ . Cf. (17) above.

The divergence of the vector product of two vectors can be easily calculated. We have

$$\begin{aligned}
 \text{div } [\mathbf{A} \cdot \mathbf{B}] &= \frac{\partial}{\partial x} [A \cdot B]_x + \frac{\partial}{\partial y} [A \cdot B]_y + \frac{\partial}{\partial z} [A \cdot B]_z \\
 &= \frac{\partial}{\partial x} (A_y B_z - A_z B_y) + \frac{\partial}{\partial y} (A_z B_x - A_x B_z) + \frac{\partial}{\partial z} (A_x B_y - A_y B_x) \\
 &= B_x \left( \frac{\partial A_z}{\partial y} - \frac{\partial A_y}{\partial z} \right) + B_y \left( \frac{\partial A_x}{\partial z} - \frac{\partial A_z}{\partial x} \right) + B_z \left( \frac{\partial A_y}{\partial x} - \frac{\partial A_x}{\partial y} \right) \\
 &\quad - A_x \left( \frac{\partial B_z}{\partial y} - \frac{\partial B_y}{\partial z} \right) - A_y \left( \frac{\partial B_x}{\partial z} - \frac{\partial B_z}{\partial x} \right) - A_z \left( \frac{\partial B_y}{\partial x} - \frac{\partial B_x}{\partial y} \right),
 \end{aligned}$$

$$\text{or} \quad \text{div } [\mathbf{A} \cdot \mathbf{B}] = (\mathbf{B} \cdot \text{curl } \mathbf{A}) - (\mathbf{A} \cdot \text{curl } \mathbf{B}). \quad . \quad . \quad (20)$$

## 14. Theorem of Coriolis.

The theory of the motion of a point relative to moving axes affords a good example of the use of vector analysis.

Consider any vector  $\mathbf{V}$  represented by the line  $OP$  in the diagram. Suppose that in time  $t$ , where  $t$  is small,  $P$  moves to  $R$  so that the vector  $\mathbf{V}$  changes from  $OP$  to  $OR$ .

Also suppose that we refer the motion of  $P$  to moving axes, and that if it remained fixed relatively to these moving axes it would move from  $P$  to  $Q$  in the time  $t$ . The displacement  $PQ$  therefore represents the motion of the axes during  $t$ . Let the motion of the axes be a rotation with angular velocity  $\omega$  about an axis  $OI$  through  $O$ . The

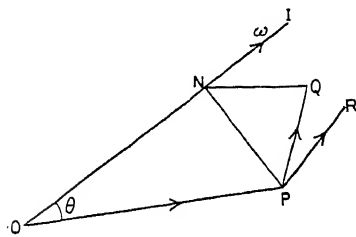


Fig. 2.

position of the axis of rotation may change with the time, but we suppose that at the instant considered it is along  $OI$ . We have then  $PQ = NP \cdot \omega t$ , where  $NP$  is the perpendicular from  $P$  to  $OI$ . But  $NP = V \sin \theta$ , so that, since  $PQ$  is perpendicular to the plane containing  $OI$  and  $OP$ , we see that

$$PQ = [\omega t \cdot \mathbf{V}].$$

The motion of  $P$  relative to the moving axes is represented by  $QR$ , and we have the vector equation

$$PR = PQ + QR,$$

or

$$PR = QR + [\omega \cdot \mathbf{V}t].$$

Now  $PR$  represents the change in  $\mathbf{V}$  in time  $t$ , so that we may put  $PR/t = \dot{\mathbf{V}}$ ; and  $QR$  represents the change in  $\mathbf{V}$  relative to the rotating axes, so that we may put  $QR/t = \dot{\mathbf{V}}'$ . Hence

$$\dot{\mathbf{V}} = \dot{\mathbf{V}}' + [\omega \cdot \mathbf{V}].$$

Here  $\dot{\mathbf{V}}$  is the rate of change of any vector  $\mathbf{V}$  relative to fixed axes, and  $\dot{\mathbf{V}}'$  the rate of change relative to axes rotating with angular velocity  $\omega$ .

Now consider the displacement  $\mathbf{D}$  of a point  $P$ , so that

$$\dot{\mathbf{D}} = \dot{\mathbf{D}}' + [\omega \cdot \mathbf{D}];$$

and also consider the velocity  $\dot{\mathbf{D}}$  of the same point, so that

$$\ddot{\mathbf{D}} = \frac{d'}{dt}(\dot{\mathbf{D}}) + [\omega \cdot \dot{\mathbf{D}}].$$

Here  $\ddot{\mathbf{D}}$  is the acceleration of the point relative to fixed axes. Hence

$$\ddot{\mathbf{D}} = \frac{d'}{dt}(\dot{\mathbf{D}}' + [\boldsymbol{\omega} \cdot \mathbf{D}]) + [\boldsymbol{\omega} \cdot (\dot{\mathbf{D}}' + [\boldsymbol{\omega} \cdot \mathbf{D}])],$$

where  $d'/dt$  indicates the rate of variation relative to the moving axes. Hence

$$\ddot{\mathbf{D}} = \ddot{\mathbf{D}}' + [\dot{\boldsymbol{\omega}}' \cdot \mathbf{D}] + [\boldsymbol{\omega} \cdot \dot{\mathbf{D}}'] + [\boldsymbol{\omega} \cdot \dot{\mathbf{D}}'] + [\boldsymbol{\omega} \cdot [\boldsymbol{\omega} \cdot \mathbf{D}]],$$

so that 
$$\ddot{\mathbf{D}} = \ddot{\mathbf{D}}' + 2[\boldsymbol{\omega} \cdot \dot{\mathbf{D}}'] + [\dot{\boldsymbol{\omega}}' \cdot \mathbf{D}] + [\boldsymbol{\omega} \cdot [\boldsymbol{\omega} \cdot \mathbf{D}]]. \quad (21)$$

This result is sometimes called *Coriolis's theorem*, after its discoverer.

If the angular velocity  $\boldsymbol{\omega}$  of the moving axes is constant in magnitude and direction,

$$\dot{\boldsymbol{\omega}}' = 0,$$

and 
$$\ddot{\mathbf{D}} = \ddot{\mathbf{D}}' + 2[\boldsymbol{\omega} \cdot \dot{\mathbf{D}}'] + [\boldsymbol{\omega} \cdot [\boldsymbol{\omega} \cdot \mathbf{D}]].$$

$[\boldsymbol{\omega} \cdot \mathbf{D}]$  is equal to  $PN \cdot \omega$ , and, since  $PN$  is perpendicular to  $\boldsymbol{\omega}$ ,  $[\boldsymbol{\omega} \cdot [\boldsymbol{\omega} \cdot \mathbf{D}]]$  is equal to  $PN \cdot \omega^2$ , and so is simply the centripetal acceleration due to the rotation. The term  $2[\boldsymbol{\omega} \cdot \dot{\mathbf{D}}']$  is called the *Coriolis acceleration*. It represents an acceleration in a plane perpendicular to the axis of rotation. If the velocity  $\dot{\mathbf{D}}'$  is directed away from the axis, in such a plane, then  $2[\boldsymbol{\omega} \cdot \dot{\mathbf{D}}']$  is equal to  $2\omega \dot{D}'$ .

Equivalent results to those above are easily found by analytical methods. For example, if the axes of  $x$  and  $y$  are rotating about  $Oz$  with any angular velocity  $\boldsymbol{\omega}$ , the velocities of the point  $P(x, y, z)$  parallel to  $Ox$  and  $Oy$  are

$$u = \dot{x} - y\omega, \quad v = \dot{y} + x\omega.$$

The accelerations of  $P$  are found from  $u$  and  $v$  in the same way as  $u$  and  $v$  are found from  $x$  and  $y$ . The component accelerations parallel to  $Ox$  and  $Oy$  are therefore

$$\dot{u} - v\omega = \ddot{x} - x\omega^2 - y\dot{\omega} - 2\dot{y}\omega,$$

and 
$$\dot{v} + u\omega = \ddot{y} - y\omega^2 + x\dot{\omega} + 2\dot{x}\omega.$$

The terms  $-2\dot{y}\omega$ ,  $2\dot{x}\omega$  are the components of the Coriolis acceleration.

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# EXAMPLES

## CHAPTER I

1. Show by means of the electromagnetic equations for empty space that in a plane electromagnetic wave the electric and magnetic fields are perpendicular to each other and to the direction of propagation of the wave.

2. By considering the electromagnetic momentum in a train of waves, show that the waves exert a pressure on a surface in which they are absorbed equal to the energy density in the waves.

3. A sphere of electricity of uniform density expands so that its radius increases while its total charge remains constant. Show that the current and the magnetic field in the sphere are everywhere equal to zero during the expansion.

4. An electron is projected with velocity  $v$  from a point at a distance  $d$  from a magnetic pole. If  $r$  denotes the distance of the electron from the pole after a time  $t$ , show that

$$r^2 = v^2 t^2 + 2av, t + d^2,$$

where  $v_r$  denotes the component of the velocity  $v$  along the direction of the magnetic field at  $t = 0$ .

5. Show that the electromagnetic angular momentum per unit length along the axis of a condenser consisting of two concentric cylinders of radii  $b$  and  $a$  is equal to  $(b^2 - a^2)he/2c$ , when the charge per centimetre on the inner cylinder is  $e$  and there is a magnetic field  $h$  along the axis.

6. Show that the force on a magnetic pole  $m$  moving with velocity  $v$  in an electric field  $F$  is equal to  $Fm(v/c)\sin\theta$ , where  $\theta$  is the angle between  $F$  and  $v$ .

7. Show that the mean rate of radiation of energy from an electron describing a simple harmonic motion of amplitude  $a$  and frequency  $\nu$  is equal to  $\frac{2}{3}(\pi/c)^3 e^2 a^2 \nu^4$ .

8. Show that

$$\frac{d}{dt} \left\{ \frac{1}{2} \int_S (\mathbf{F}^2 + \mathbf{H}^2) dS \right\} + \int_S \rho (\mathbf{F} \cdot \mathbf{V}) dS + c \int_\sigma [\mathbf{F} \cdot \mathbf{H}]_n d\sigma = 0,$$

where  $\mathbf{F}$  and  $\mathbf{H}$  are the electric and magnetic field strengths,  $\rho$  the density of electricity,  $\mathbf{V}$  the velocity of the electricity,  $S$  is the volume enclosed by a surface of area  $\sigma$ , and the suffix  $n$  indicates the component along the normal to the element of area  $d\sigma$ .

9. A condenser consisting of two concentric cylinders with radii  $b$  and  $a$ , with the space between them filled with a medium of specific inductive capacity  $K$  and magnetic permeability  $\mu$ , is rotating with angular velocity  $\omega$  about its axis in a magnetic field  $H$  parallel to its axis. Show that if the cylinders are connected together there are equal and opposite charges on them equal to  $\pm \pi\omega H(\mu K - 1)(b^2 - a^2)/c \log(b/a)$  per unit length along the axis.

10. A hollow circular cylinder of specific inductive capacity  $K$  is placed in a radial electric field of strength  $A/r$  at a distance  $r$  from the axis of the cylinder. If the cylinder is rotating with angular velocity  $\omega$ , calculate the magnetic field in the cylinder.

## CHAPTER II

1. Calculate the magnetic field at a point on the axis of a circle round which an electron is moving with uniform velocity. Show that its component along the axis is equal to the field due to a current, round the circle, equal to the charge on the electron multiplied by the number of revolutions it makes per second.

2. A magnetic field  $H$  is suddenly generated perpendicular to the plane of a uniform circular ring of electricity of charge  $e$ , mass  $m$ , and radius  $a$ . Calculate the resulting angular velocity of the ring, assuming the field due to the ring very small compared with  $H$ .

3. What would be the apparent magnetic permeability of a long uniform bar the material of which was a perfect conductor of electricity?

4. In the case of a large number of similar systems each having possible energies  $0, \epsilon, 2\epsilon, 3\epsilon, \dots, n\epsilon, \dots$ , show that in the equilibrium state the fraction for which the energy  $E$  is between  $E$  and  $E + dE$  is equal to  $\epsilon^{-E/kT} dE/kT$  when  $\epsilon$  is made indefinitely small.

5. If  $\mathbf{I}$  is defined as the magnetic moment per unit volume, and  $\mathbf{B} = \mathbf{H} + 4\pi\mathbf{I}$ , where  $\mathbf{H}$  is the magnetic field strength, defined as usual as the field inside a long narrow cavity parallel to  $\mathbf{I}$ , show that  $\text{div}\mathbf{B} = 0$ .

6. A space in which there is a uniform magnetic field  $H$  contains  $n$  electrons per cubic centimetre all moving with velocities of the same magnitude  $v$  in planes perpendicular to  $H$ . Neglecting the mutual action of the electrons, calculate the intensity of magnetization. Ans.  $\frac{1}{2}nmv^2/H$ .

7. Discuss the analogy between the "critical temperature" at which ferromagnetic properties disappear and the "critical temperature" above which a vapour cannot be liquefied.

8. Calculate the magnetic field at the centre of the circular electron orbit of a Bohr magneton.

9. An electron of mass  $m$ , spinning so that it has a magnetic moment equal to that of a Bohr magneton, is projected with velocity  $v$  from a point at a distance  $d$  from a north pole  $P$ . If  $v_r$  is its initial velocity along the direction of the field due to the pole, and if its magnetic axis points towards the pole, how far from the pole will it get? Ans.  $\sqrt{[(9.2 \times 10^{-21}P)/(9.2 \times 10^{-21}P/d^2 - \frac{1}{2}mv_r^2)]}$ .

10. In Stern and Gerlach's experiment, if  $l = 5$  cm. and  $\partial H/\partial y = 100,000$ , calculate the mean deviation  $y$  for silver atoms having a temperature of  $1000^\circ \text{C}$ .

## CHAPTER III

1. Show that, on the classical theory of thermionic emission, the mean kinetic energy of the electrons which escape from the metal is equal to that of the electrons inside the metal as they enter the surface layer.

2. The thermionic current  $i$  from a wire is found to be represented by the equation  $i = AT^2\epsilon^{-b/T}$ , where  $A$  and  $b$  are constants. Show that it can be equally well represented by  $i = A'T^2\epsilon^{-b'/T}$ , where  $A'$  and  $b'$  are not equal to  $A$  and  $b$ , provided that

$$T \log \frac{A}{A'} = b - b'.$$

3. If the latent heat of evaporation of the electrons from a metal is given by

$L = \frac{RT^2}{p} \frac{dp}{dT}$ , where  $p$  is the pressure of the electron gas, and if  $L = L_0 + aT$ , where  $a$  is a constant, show that the thermionic current density is given by  $i = AT^{a/R - \frac{1}{2}} e^{-L_0/RT}$ .

4. Calculate the maximum thermionic current from a straight wire, 0.1 mm. in diameter and 10 cm. long, to a cylinder surrounding it, of 2 cm. radius, due to a potential difference of 100 volts.

5. What fraction of the saturation thermionic current from a wire at 1500° C. would be obtained with an electrode surrounding the wire at a potential of one volt above that of the wire?

## CHAPTER IV

1. If a plane surface, at potential  $V$ , illuminated by ultra-violet light, emits electrons, equally in all directions and all with velocity  $v$ , from every point illuminated, show that the current to a parallel plane at potential zero is equal to  $ne[1 - \sqrt{(2Ve)/(mv^2)}]$ , where  $n$  is the total number of electrons emitted per second.

2. The potential difference between two large parallel metal plates is  $V$ . There is a uniform magnetic field  $H$  in the space between them parallel to the surfaces of the plates. A small area on the negative plate is illuminated and emits  $n$  electrons per second with negligible velocities. Calculate the current between the plates when  $n$  is so small that the space charge can be neglected.

3. If  $\frac{1}{2}c \int_0^\infty T_\nu F(\nu) E(\nu) d\nu = 2NkT$  (p. 64), show that  $T_\nu = h\nu - w_0$ .

4. Two plane circular metal plates A and B, each 25 cm. in diameter, are supported 2 cm. apart. The potential difference between the plate A and the ground is indicated by a quadrant electrometer which gives a deflection of 1000 mm. for one volt. If the potential of the plate B is changed from 0 to 1 volt, the electrometer deflection changes 200 mm. If the plate B is kept at -100 volts, A being at zero potential, and B is illuminated for 10 sec. by ultra-violet light, the electrometer gives a deflection of 100 mm. Calculate the current between the plates due to the light. Also calculate the apparent capacity of the electrometer. Ans. Current,  $1.08 \times 10^{-12}$  amp., capacity 78 cm.

5. How many quanta per second of frequency  $\nu = 3 \times 10^{14}$  per second falling normally on a black plate would give a force on the plate of 1 dyne? If the plate had a mass of 1 gm. and specific heat 0.1 calorie per gramme per degree Centigrade, at what rate would its temperature rise? Ans.  $1.53 \times 10^{22}$ ;  $7.16 \times 10^3$  degrees Centigrade per second.

## CHAPTER V

1. Show that the number of ways in which  $N$  like objects can be arranged in  $n$  boxes is equal to  $n^N$ . Hence show that when a gas containing  $N$  atoms expands freely from a volume  $V_1$  to a volume  $V_2$  its entropy is increased by  $kN \log(V_2/V_1)$ .

2. The energy of a collection of  $\mathcal{N}$  like systems, each of which can have energy 0 or  $\epsilon$  only, is equal to  $\frac{1}{2}\mathcal{N}\epsilon$ . Calculate the temperature of the collection.

3. If  $N$  objects are distributed at random among  $M$  boxes, show that, when  $M$  and  $N$  are very large, the fraction of the boxes which get  $n$  objects is equal to  $\frac{M}{N+M} \left( \frac{N}{N+M} \right)^n$ . Show that this result agrees with the equilibrium distribution of energy among  $\mathcal{N}$  systems, each of which can have energies 0,  $\epsilon$ ,  $2\epsilon$ ,  $3\epsilon$ , ...

4. Deduce Maxwell's law for the distribution of the velocities among the molecules of a gas from the quantum theory of a monatomic gas.

5. Calculate the vapour pressure of mercury at  $0^\circ \text{C}$ ., in millimetres of mercury, by means of the equation (p. 75)

$$\log p = -\frac{L}{k\mathcal{N}\theta} + \log_e \left\{ \left( \frac{2\pi m}{h^2} \right)^{\frac{3}{2}} (k\theta)^{\frac{5}{2}} \right\}.$$

6. Show that, according to Debye's theory of specific heats, the shortest possible wave-length in a solid is nearly equal to the mean distance between the centres of two adjacent atoms.

7. Assuming that heat radiation consists of quanta of energy  $h\nu$  and that the pressure  $p$  on the walls of a vessel containing radiation of energy density  $E$  is equal to  $E/3$ , show that  $p = 0.90 \dots \mathcal{N}k\theta$ , where  $\mathcal{N}$  is the number of quanta per cubic centimetre,  $k$  the gas constant for 1 molecule, and  $\theta$  the absolute temperature.

8. If an electron of mass  $m$  and charge  $e$  is describing a circular orbit about a fixed positive charge  $E$ , and if the angular momentum of the electron can only be equal to  $n\hbar/2\pi$ , where  $n = 1, 2, 3, 4, \dots$ , show that the energy is equal to  $A - (2\pi^2 e^2 E^2 m)/(n^2 \hbar^2)$ , where  $A$  is a constant.

9. If the acceleration of a particle relative to fixed axes is  $\mathbf{A}$ , show that

$$\mathbf{A} = \frac{d^2 \mathbf{R}}{dt^2} + 2 \left[ \boldsymbol{\omega} \cdot \frac{d\mathbf{R}}{dt} \right] + \left[ \boldsymbol{\omega} \cdot [\boldsymbol{\omega} \cdot \mathbf{R}] \right],$$

where  $\mathbf{R}$  is the distance of the particle from the origin relative to axes rotating with constant angular velocity  $\boldsymbol{\omega}$ .

10. For a pendulum consisting of a particle of mass  $m$ , suspended by a thread of length  $l$ , describing a circle in a horizontal plane, show that according to the quantum theory the possible values of  $r$ , the radius of the circle, are given by

$$\frac{r^3}{l^2 - r^2} = \frac{n^4 \hbar^4}{16\pi^4 m^4 g^2},$$

where  $n = 0, 1, 2, 3, \dots$ . Show also that if  $l$  is slowly made shorter the kinetic energy divided by the frequency will remain constant.

11. For a particle of mass  $m$  moving along the  $x$  axis and acted on by a force equal to  $(-\mu x)$ , Schrödinger's equation  $\Delta\psi + \frac{8\pi^2 m}{h^2} (h\nu - V)\psi = 0$  becomes

$$\frac{\partial^2 \psi}{\partial x^2} + \frac{8\pi^2 m}{h^2} (h\nu - \frac{1}{2}\mu x^2)\psi = 0. \text{ Transform this into } \frac{\partial^2 \psi}{\partial q^2} + (C - q^2)\psi = 0, \text{ where}$$

$$C = 2\nu/\nu_0 \text{ and } \nu_0 = \frac{1}{2\pi} \sqrt{\frac{\mu}{m}}. \text{ Find a solution in the form } \psi = e^{-\frac{1}{2}q^2} \sum_0^\infty a_n q^n, \text{ and}$$

show that if  $C = 2n + 1$  ( $n = 0, 1, 2, 3, \dots$ ) the series terminates.

## CHAPTER VI

1. Calculate the potential difference in volts through which an electron must drop in order to have enough energy to move the electron in a hydrogen atom from the orbit with quantum number 2 to that with number 3.

2. Calculate the average kinetic energy in ergs of electrons from a wire at  $2000^\circ \text{C}$ . What potential difference in volts would be required to stop an electron having this average energy?

3. The potential difference between two parallel plates is equal to  $V$ . The space between them contains a gas at a low pressure, the molecules of which are ionized by electrons which have fallen through  $A$  volts, but collisions with slower electrons are perfectly elastic. If the negative plate emits  $n$  electrons, show that

the charge received by the positive plate is equal to  $ne \cdot 2p$ , where  $p$  is equal to the integral part of  $V/4$ , assuming there is no appreciable recombination.

4. Calculate the ionization potential of an atom consisting of a heavy sphere of electricity of uniform density, charge  $e$ , and radius  $a$ , with an electron with charge  $-e$  at the centre. What value of  $a$  would make the ionization potential  $V$  agree with that given by the equation  $Ve = h\nu$ , where  $\nu$  is the frequency of the electron?

5. Calculate the ionization potentials of a helium atom, assuming it to consist of a heavy nucleus with charge  $2e$  and two electrons moving round a circle at opposite ends of a diameter, the angular momentum being equal to  $h/\pi$ . When the atom has lost one electron, assume that the other one moves in a circle with angular momentum  $h/2\pi$ .

## CHAPTER VII

1. An X-ray tube uses 30 milliamperes at 100,000 volts. Assuming that 1 per cent of the energy goes into X-rays which are emitted equally in all directions over a hemisphere, calculate the energy of the rays falling on 1 sq. cm. per second at a distance of 1 m. from the anti-cathode.

2. Calculate the wave-length of X-rays for which the energy of one quantum is  $10^{-6}$  erg.

3. A crystal consists of equal atoms arranged on a cubical space lattice, the cubical elements having sides  $5 \times 10^{-8}$  cm. long. Calculate the first order glancing angles for X-rays of wave-length  $10^{-8}$  cm. from the (100), (210), and (112) faces.

4. The refractive index  $\mu$  of a substance for light of frequency  $\nu$  is given by

$$\mu^2 = 1 + \frac{1}{2} \frac{\nu_s^2}{\nu_s^2 - \nu^2},$$

where  $\nu_s = 3 \times 10^{13}$ . Calculate the refractive index for X-rays of wave-length  $10^{-8}$  cm., and the critical glancing angle for total reflection of these rays at the surface of the substance.

5. If X-rays are reflected from the surface of a crystal the refractive index of which for the rays is  $\mu$  and grating space  $d$ , show that

$$n\lambda = 2d\mu \sqrt{1 - (1/\mu) \cos^2 \theta}.$$

6. If  $\theta_1$  is the value of  $\theta$  found when  $n = n_1$ , in the previous example, and  $\theta_2$  that with  $n_2$ , show that

$$1 - \mu = \frac{n_2^2 \sin^2 \theta_1 - n_1^2 \sin^2 \theta_2}{n_2^2 - n_1^2}.$$

Putting  $n_1 \lambda_1 = 2d \sin \theta_1$ , and  $n_2 \lambda_2 = 2d \sin \theta_2$ , and  $\lambda_1 - \lambda_2 = \delta \lambda$ , show that when  $\delta \lambda / \lambda_1$  is small,

$$1 - \mu = \frac{2n_2^2 \sin^2 \theta_1 \delta \lambda}{(n_2^2 - n_1^2) \lambda}.$$

7. Electrons are projected with velocity  $v$  from a point in a uniform magnetic field  $H$ . The electrons all start off in nearly the same direction, which is perpendicular to the field. Show that they will all pass through a line parallel to the field, and find the position of this line.

8. Show that according to the classical theory the energy scattered per second by an electron, when in a beam of plane polarized X-rays, inside a cone of solid angle  $d\omega$  making an angle  $\theta$  with the direction of the electric field in the rays, is equal to  $(\bar{F}^2 e^4 \sin^2 \theta d\omega) / (4\pi m^2 c^3)$ , where  $\bar{F}$  denotes the root mean square of the electric field in the incident rays.

Hence show that for unpolarized rays the energy per second in a cone making an angle  $\varphi$  with the direction of the rays is  $(\bar{F}^2 e^4 / 8\pi m c^3) (1 + \cos^2 \varphi) d\omega$ .

9. Show that it is impossible for a free electron initially at rest to completely absorb a quantum of radiation.

10. Apply A. H. Compton's theory of the scattering of X-rays to the case of a quantum  $h\nu$  being scattered by an electron moving initially with velocity  $v$  in a direction making an angle  $\theta$  with that in which the quantum is moving.

11. Calculate the change in wave-length when light of wave-length  $10^{-4}$  cm. is scattered by free electrons in a direction perpendicular to the direction of propagation of the light. Would you expect to get this change of wave-length when light is reflected from a metallic mirror?

12. Show that the change of wave-length due to scattering of X-rays by free electrons on the quantum theory is the same as that indicated by the classical theory for scattering by electrons moving in the direction of the incident rays with a certain velocity, and calculate this velocity.

## CHAPTER VIII

1. The wave numbers of a spectral series are 4857, 19,932, 25,215, 27,665, 28,997. Show that the limit of this series is approximately 32,033.

2. The value of the constant  $N$  in the formula

$$\bar{\nu} = Z^2 N \left( \frac{1}{n^2} - \frac{1}{m^2} \right)$$

for hydrogen is 109,678.3 and for helium 109,722.9. Show that for an atom with a very large value of  $Z$  it is 109,737.7.

3. Could lines of wave numbers 3427.5, 3808.3 be lines in the spectrum of atomic hydrogen?

4. The wave numbers in a band spectrum are given by  $\nu = 20,000 \pm 1000n + 90n^2$ . Calculate the wave number for the head of the band.

5. The wave numbers in a band spectrum are given by  $\nu = 1000(2n - 1)$  when  $n$  is positive, and  $\nu = -1000(2n + 1)$  when  $n$  is negative. Calculate the moment of inertia of the molecules emitting this spectrum.

## CHAPTER IX

1. A light flexible wire lies in a plane which is perpendicular to a non-uniform magnetic field. If the tension in the wire is  $T$ , and it carries a current  $c$ , show that it lies along a possible path of an electron in the field, provided

$$\frac{T}{c} = \frac{mv}{e},$$

where  $m$  is the mass,  $e$  the charge, and  $v$  the velocity of the electron.

2. A particle of mass  $m$  and charge  $e$  is projected from the origin of co-ordinates  $x, y, z$  along the  $x$  axis with velocity  $v$ . There is a magnetic field  $H$  along the  $z$  axis. If  $y_l$  is the  $y$  co-ordinate of a point on the path of the particle at  $x = l$ , show that provided  $y/x$  is small

$$y_l = \frac{e}{mv} \int_0^l (l - x) H dx.$$

3. If instead of the magnetic field, in the previous example, there is an electric field  $Y$  along the  $y$  axis, show that

$$y_1 = \frac{e}{mv^2} \int_0^l \left( \int_0^x Y dx \right) dx.$$

4. Show that the equations of motion of a particle of mass  $m$  and charge  $e$  in a magnetic field with  $x, y, z$  components  $F, G, H$  are  $m\ddot{x} = e(H\dot{y} - G\dot{z})$ ,  $m\ddot{y} = e(F\dot{z} - H\dot{x})$ ,  $m\ddot{z} = e(G\dot{x} - F\dot{y})$ , and show by means of these equations that the velocity of the particle remains constant.

5. If  $F$  and  $G$  are zero and there is an electric field  $Y$  along the  $y$  axis, show that

$$m\ddot{x} = eH\dot{y}, \quad m\ddot{y} = -eH\dot{x} + Ye, \quad m\ddot{z} = 0.$$

Show that a solution is given by  $\dot{x} = A \sin(\omega t + \alpha) + Y/H$ ,  $\dot{y} = A \cos(\omega t + \alpha)$ ; and find the significance of the constants  $A, \omega, \alpha$ .

6. A particle carrying a charge  $e$  and having initial kinetic energy  $T$  passes through the radial electric field between two concentric spheres of which the outer has radius  $R$  and the inner a charge  $E$ . Show that

$$h = \left\{ \frac{1}{R^2} + \left( \frac{1}{k} - \frac{1}{R} \right)^2 \tan^2 \left( \frac{\phi}{2} \right) \right\}^{-\frac{1}{2}},$$

where  $h$  is the perpendicular from the centre of the spheres to the original direction of the path,  $k = -Ee/2T$ , and  $\phi$  is the angle through which the path is deflected.

7. The velocity  $v$  of an  $\alpha$ -ray which has travelled a distance  $x$  is given by  $v^2 = A(R-x)$ . Show that the time it takes to go a distance  $x$  is  $t = \frac{2}{3} A^{-\frac{1}{2}} \{ R^{\frac{3}{2}} - (R-x)^{\frac{3}{2}} \}$ . Calculate the force on the particle as a function of  $x$ , and as a function of  $t$ .

8. If an  $\alpha$ -ray makes a head-on collision with a hydrogen atom at rest, and the collision is perfectly elastic, calculate the ratio of the velocity given to the hydrogen atom to the initial velocity of the  $\alpha$ -ray.

9. If  $\alpha$ -rays are scattered by a sheet of gold containing  $10^{15}$  atoms per square centimetre, and the initial velocity of the rays is  $2 \times 10^9$  cm. per second, what fraction of the rays will be scattered through angles between  $80^\circ$  and  $100^\circ$ ?

10. Calculate the potential difference in volts required to give an  $\alpha$ -ray a velocity of  $2 \times 10^9$  cm./sec.

## CHAPTER X

1. What magnetic and electric fields would be necessary in J. J. Thomson's positive ray parabola apparatus (p. 188) to give equal electric and magnetic deflections of 2 cm. with positive rays consisting of oxygen atoms carrying the protonic charge  $e$ . Take  $d = 5$  cm.,  $l = 20$  cm., and  $v = 10^9$  cm./sec.

2. Mention several possible positive rays for which  $e/m = 603$  approximately.

3. In Aston's mass spectrograph (fig. 4, p. 191), if the angle between AB produced and BD is denoted by  $\theta_0$  and is supposed constant, show that for BD to be independent of the velocity of the rays we must have  $\theta = \theta_0$ .

4. In Aston's mass spectrograph (fig. 4, p. 191) show that if  $y = BD$ , then  $dy/y = -\theta(m/e)d(e/m)$ , approximately.

5. In Dempster's method of finding  $e/m$  (p. 193), show that  $dr/r = -\frac{1}{2}(m/e)d(e/m)$ .

## CHAPTER XI

1. Show that the average life of the atoms of a radioactive body which decays according to the equation  $N_t = N_0 e^{-\alpha t}$  is equal to  $1/\alpha$ .

2. Show that the average life of the atoms of a radioactive body which disappear during an interval  $t$  is equal to

$$\frac{1 - (1 + \alpha t)e^{-\alpha t}}{\alpha(1 - e^{-\alpha t})}.$$

When  $t$  is very small, show that this reduces to  $\frac{1}{2}t$ .

3. Calculate the weight in grammes of 1 c. m. of radon gas at  $0^\circ\text{C}$ . and 760 mm.

4. One milligramme of radium is in a small cavity of radius 0.5 mm. at the centre of a sphere of radius 2 cm. and thermal conductivity 0.01. If the surface of the sphere is kept at  $0^\circ\text{C}$ ., what will be the temperature of the radium?

5. After how many years would initially pure uranium contain 1 per cent by weight of lead?

## CHAPTER XII

1. Do Millikan's experiments show that the charge on an electron is exactly equal, though of opposite sign, to the charge on a proton? Does this follow from the fact that bodies containing enormous numbers of electrons and protons can be obtained without any sensible charge on them?

2. It has been proposed to call an electrically neutral particle formed by the combination of a proton and an electron a neutron. Why do not all the protons and electrons neutralize each other, forming neutrons? What should we expect the properties of a substance composed of neutrons to be?

3. It is found that the physical properties of nitrogen and carbon monoxide are very similar. How can this be explained?

4. If an  $\alpha$ -ray consists of 6 point charges  $\pm e$  arranged at equal distances apart along a straight line, with two positive charges in the middle and one at each end, show that the energy lost in the formation of the  $\alpha$ -ray is approximately  $2.752e^2/a$ , where  $a$  is the distance between adjacent charges. Hence calculate  $a$  by assuming this energy lost equal to that required to explain the difference between four times the atomic weight of hydrogen and that of helium.

5. If matter consists of nothing but electric point charges attracting and repelling according to the inverse square of the distance, show that there is nothing to determine a definite size for any piece of matter or any atom composing the piece.

## CHAPTER XIII

1. The gas between two parallel plates at a distance  $d$  apart is ionized so that  $q$  positive and  $q$  negative ions are produced per cubic centimetre per second. Show that the maximum or saturation current density is equal to  $dqe = I$ . If the current density  $i$  is less than  $I$ , show that, assuming  $n_1 = n_2$  everywhere,

$$I - i = \frac{\alpha d^3 i^2}{P^2 e(k_1 + k_2)^2}$$

approximately, where  $P$  is the potential difference between the plates and  $\alpha$  is the coefficient of recombination.

2. In Zeleny's method of finding  $k$  (p. 215), show that, if  $b - a = h$  is a small quantity, then

$$k = \frac{vh}{Xd},$$

where  $v$  is the average velocity of the gas and  $X$  the electric field strength.

3. If  $v$  is the velocity of the gas between two parallel plates at a distance  $d$  apart, and ions start from one plate and reach the other at a distance  $h$ , in the direction of  $v$ , from the starting-point, show that

$$h = \frac{1}{k} \int_0^d \frac{v dx}{X},$$

where  $X$  is the electric field at a distance  $x$  from the plate at which the ions start.

4. In Langevin's method of finding  $k$  (p. 216), if instead of ionizing the gas over the whole distance between the plates it is only ionized in a thin layer midway between the plates, show how the charge received by the plates will vary with the time interval  $t$ .

5. Assuming that an ion moving through a gas with velocity  $v$  is retarded by a force equal to  $(-\mu v)$ , show that the velocity of the ion due to an electric field  $X$  is equal to  $Xe\delta/mW$ , where  $\delta$  is the distance the ion moves through the gas when projected with initial velocity  $W$ .

6. According to the kinetic theory of gases the coefficient of diffusion is approximately given by  $K = \frac{1}{3}\lambda V$ , and the velocity of an ion due to a field  $X$  is  $kX = Xe\lambda/mF$ . Deduce the equation  $k = K\mathcal{N}e/RT$  (p. 220) from these equations.

7. Calculate the total number of electrons in a drop of water which falls in air at the rate of 1 mm. per second.

8. Is it possible for a charged drop of mercury which falls in air at the rate of 1 mm. per second to remain at rest in a vertical field of 5000 volts per centimetre?

9. A number  $N$  of molecules in a gas is uniformly distributed over the volume of a sphere of radius  $a$ . Calculate the mean square of the distance of these molecules from the centre of the sphere after a time  $t$ .

10. Show that if  $N$  molecules in a gas describe  $n$  free paths, the mean square of the distance of these molecules from any point is increased by  $2n\lambda^2$ , where  $\lambda$  is the mean free path.

11. The vapour pressure  $P$  of a spherical uncharged drop of radius  $a$  is greater than that of a plane surface by

$$\delta P = \frac{\rho'}{\rho} \frac{2T}{a},$$

where  $\rho'$  is the density of the vapour,  $\rho$  that of the liquid, and  $T$  the surface tension of the liquid. If the drop carries a charge  $E$ , show that

$$\delta P = \frac{\rho'}{\rho} \left( \frac{2T}{a} - \frac{E^2}{8\pi a^4} \right).$$

Hence calculate the radius of a water drop carrying the protonic charge  $e$  for which  $\delta P = 0$ . The surface tension of water is 80 dynes/cm.

12. Air saturated with water vapour at 16° C. and 760 mm. is suddenly expanded in the ratio 1:1.5. Calculate the amount of water which condenses, and the final temperature.

## CHAPTER XIV

1. In Townsend's apparatus (p. 236), if the slit at S is replaced by a small hole at the centre of the disc B, and the lower plate GHK is divided into a circular disc of radius  $a$  surrounded by a concentric ring, show that the fraction of the charge reaching the disc of radius  $a$  is  $1 - e^{-\alpha a^2}$ , where  $\alpha = Ze/4\pi kTza$ , and  $za$  is the value of  $z$  at the lower plate.

2. Assuming atoms of argon to be hard spheres and electrons to be points, calculate the diameter of an argon atom if the mean free path of a high-velocity electron in argon at 1 mm. pressure is 0.147 cm. Ans.  $7.8 \times 10^{-9}$  cm.

3. Show that when an electron is moving in a strong electric field  $X$  in a gas at pressure  $p$ , and we suppose that the velocity of the electron is reduced to a small value by each collision, then its velocity through the gas is proportional to  $\sqrt{X/p}$ .

4. If  $\alpha$  denotes the number of pairs of ions produced by an electron per centimetre as it moves in a field  $X$  in a gas at pressure  $p$ , and if we suppose that the electron moves in a straight line, show that  $\alpha = (p/\lambda_1)e^{-pV/X\lambda_1}$ , where  $\lambda_1$  is the free path of the electron at 1 mm. pressure, and  $V$  is the potential through which the electron must drop to get enough energy to ionize a molecule. Assume that every collision at which the energy of the electron is greater than  $V_e$  gives ionization and that at every collision the velocity of the electron is reduced to zero.

5. If the electric field between two parallel plates is not uniform, owing to the presence of unequal numbers of ions and electrons, show that, assuming the electrons produce ions by collisions but that the ions do not,

$$\int_0^D \alpha e^{-\int_0^x \alpha dx} dx = \frac{1}{1+\gamma}$$

is the condition for a continuous discharge, where  $\gamma$  is the number of electrons liberated at the cathode by the impact of a positive ion,  $\alpha$  is the number of pairs of ions produced by an electron per centimetre, and  $D$  is the distance between the plates. If  $X$  is constant, show that the above condition reduces to  $e^{-\alpha D} = \gamma/(1+\gamma)$ . If  $\gamma$  is a constant, and  $\alpha = pf(X/p)$ , show that the condition for the potential difference to be a minimum is  $pD = Vf'(V/pD)/f(V/pD)$ .

## CHAPTER XV

1. In the theory of the variation of the potential between parallel electrodes, with an ionized gas between them, show that if, in the layer of thickness  $\lambda_1$ , at the positive electrode, we assume  $q - \alpha n_1 n_2 = q(1 - x/\lambda_1)$ , then the potential drop  $V_1$  at this electrode is given by  $V_1^2 = \frac{32}{75} \pi i \lambda_1^3 / k_2$ .

2. The current in amperes through a flame between two parallel electrodes at a distance  $d$  cm. apart was given by the equation

$$V = 3 \times 10^6 C d + 6 \times 10^{14} C^2,$$

where  $C$  = current and  $V$  = potential difference. The cross-section of the flame was about 4 sq. cm. If the velocity of the negative ions due to a field of 1 volt per centimetre was 7000 cm./sec., calculate the number of negative ions per cubic centimetre in the flame. Ans.  $7.5 \times 10^7$ .

3. Calculate the thickness of the layer in which the field varies, at the cathode, in the flame of the previous example, taking the mobility of the positive ions to be 2 cm./sec. per volt/centimetre, when the current is  $10^{-6}$  amp. and  $d = 10$  cm. Ans.  $\lambda_2 = 0.63$  cm.

4. If one electrode in the flame of example 2 is coated with lime or potassium carbonate, show that the current due to 630 volts will be 210 times greater in one direction than in the other.

5. Show that  $\log_{10} K_2/K_1 = 5048(V_1 - V_2)/T$ , where  $K_2$  and  $K_1$  are the equilibrium constants for the equilibrium between the vapours of two metals and electrons at the absolute temperature  $T$ , and  $V_1$  and  $V_2$  are the ionization potentials of the metals in volts.

6. The ionization potentials of caesium, rubidium, potassium, sodium, and lithium are 3.87, 4.15, 4.32, 5.11, and 5.36 volts respectively. Show that the relative numbers of atoms of these metals required to give equal conductivities to a flame at 2000° C. are approximately as 1, 4.8, 13.2, 1330, 5731, when the amounts are not very small. (The relative numbers of atoms to give equal conductivities to a Bunsen flame are found experimentally to be about as 1, 4.5, 7.3, 630, 4400.)

## CHAPTER XVI

1. In a uniform positive column, assuming all the ionization to be due to collisions by the electrons, and all the loss of ions due to diffusion, and the rate of loss to be inversely as the pressure and proportional to the number of electrons per cubic centimetre, show that the relation between the electric field  $X$  and the pressure  $p$  must be of the form  $p^2 f(X/p) = \text{constant}$ , where  $f(X/p)$  denotes a function of  $X/p$ . If  $X \propto \sqrt{p}$ , find  $f(X/p)$ .

2. If in an electric discharge through a gas the electric intensity  $X$  diminishes as the current density increases, show that the cross-section of the discharge will contract until the current density has the value for which  $X$  is a minimum.

3. The "normal" cathode fall of potential is approximately equal to the minimum sparking potential. Discuss this result.

4. The variation of the potential in a discharge tube close to the anode is similar to that near the cathode but on a much smaller scale. Why is this?

5. It has been found that a uniform positive column in a discharge tube, in a magnetic field perpendicular to its length, moves sideways in a direction perpendicular to the magnetic field and to the length of the column, with a velocity approximately proportional to the magnetic field  $H$  and inversely to the pressure  $p$ . Show that we should expect the velocity to be equal to  $Hv_1v_2/X$ , where  $v_1$  and  $v_2$  are the velocities of the positive and negative ions along the column, and  $X$  is the electric intensity.

## CHAPTER XVII

1. If 34 pairs of ions are generated per cubic centimetre per second in the air, how many positive ions will there be per cubic centimetre? Coefficient of recombination =  $3400e$ , with  $e$  in electrostatic units. Ans. 4600.

2. What will be the conductivity of the air in the previous example, in electrostatic units, if  $k_1 = k_2 = 1.6 \text{ cm./sec. per volt/cm.}$  Ans. 0.0021.

3. If 3 cm. of rain fall, consisting of drops 0.2 cm. in diameter, each carrying a charge of  $4.77 \times 10^{-10}$  electrostatic units, what will be the charge received by the earth per square kilometre. Ans. 456 e.s.u.

4. A cloud consisting of a sphere of radius 0.5 km., with its centre 4 km. above the earth's surface, discharges to the earth. If there are  $8.4 \times 10^4$  drops per cubic centimetre in the cloud, each carrying a charge  $4.77 \times 10^{-10}$  e.s.u., what will be the change in the vertical electric field on the earth's surface, at a point 10 km. from the point vertically below the centre of the cloud? Ans. 4 volts/cm.

5. If there were a current of 500 amp. into the earth from the air in the northern hemisphere, and a current of 500 amp. from the earth into the air in the southern hemisphere, how much work in ergs would be required to take a unit magnetic pole once round the equator? Ans. 630.

## CHAPTER XVIII

1. Show that  $x^2 + y^2 + z^2 - c^2 t^2$  is transformed into  $x'^2 + y'^2 + z'^2 - c^2 t'^2$  by the transformation

$$x = x' \cosh \alpha + ct' \sinh \alpha, \quad y = y',$$

$$t = t' \cosh \alpha + \frac{x'}{c} \sinh \alpha, \quad z = z'.$$

If  $\alpha = \frac{1}{2} \log \left( \frac{c+v}{c-v} \right)$ , show that this transformation agrees with that of the special theory of relativity.

2. If the electric field in a ray of light travelling along the  $x$  axis is proportional to  $\sin \frac{2\pi}{\lambda} (x - ct)$  to an observer at rest relative to the axes  $x, y, z$ , show that to an observer moving along the  $x$  axis with velocity  $v$  the field in the ray will be proportional to

$$\sin \frac{2\pi}{\lambda'} (x' - ct'),$$

where

$$\frac{\lambda}{\lambda'} = \frac{1 - v/c}{\sqrt{1 - v^2/c^2}}.$$

3. If an observer is moving along the  $x$  axis with a velocity  $v$  and observes the volume of a body moving along the  $x$  axis with a velocity  $u$ , show that the ratio of the volume observed to that of the body when  $u = v$  is

$$\sqrt{\frac{(c^2 - u^2)(c^2 - v^2)}{(c^2 - uv)^2}}.$$

4. Considering only two co-ordinates  $x$  and  $t$ , and taking the velocity of light  $c = 1$ , show that changing from rectangular axes  $x$  and  $t$  to new ones, which are moving relatively to  $x$  and  $t$  with velocity  $v$  along the  $x$  axis, is equivalent to changing to oblique axes with the new  $x$  axis,  $x'$ , making an angle  $\psi$  with the old  $x$  axis and the new  $t$  axis,  $t'$ , making an angle  $\psi$  with the old  $t$  axis so that the angle between  $t'$  and  $x'$  is  $\frac{1}{2}\pi - 2\psi$ . Show that  $\tan \psi = v$ .

5. In example 4, show that the intercepts of the curves  $x^2 - t^2 = \pm 1$  on the axes  $x'$  and  $t'$  are equal to the units of length and time in the moving rectangular axes.

6. Show that, if  $H_x, H_y, H_z$  and  $F_x, F_y, F_z$  are the components of an electromagnetic field relative to axes  $x, y, z$ , the components relative to axes  $x', y', z'$  moving along  $x$  with velocity  $v$  and coinciding with  $x, y, z$  at  $t = 0$  are given by

$$H_{x'} = H_x, \quad H_{y'} = \beta \left( H_y + \frac{v}{c} E_z \right), \quad H_{z'} = \beta \left( H_z - \frac{v}{c} E_y \right),$$

$$E_{x'} = E_x, \quad E_{y'} = \beta \left( E_y - \frac{v}{c} H_z \right), \quad E_{z'} = \beta \left( E_z + \frac{v}{c} H_y \right),$$

$$\text{where } \beta = \frac{1}{\sqrt{1 - v^2/c^2}}.$$

7. A body of mass  $M$  moving through empty space, with a velocity  $V$ , along a straight line, loses particles of matter, as it goes along, in such a way that the particles remain at rest. Show that  $\frac{d}{dt}(MV) = 0$ , so that it may be said that there is no force on the body, since its momentum remains constant although its velocity increases as  $M$  diminishes. If, however, the particles continue to move with velocity components, parallel to the path of the body, equal to  $V$ , show that  $dV/dt = 0$ , so that the momentum of the body diminishes at the rate  $(-V \frac{dM}{dt})$ , and there may therefore be said to be a retarding force acting on the body equal to the rate of change of its momentum. Criticize these statements.

8. Show that the difference  $w$  between two velocities  $v$  and  $u$ , as defined in the special theory of relativity, is given by the equation

$$\tanh^{-1} \frac{w}{c} = \tanh^{-1} \frac{v}{c} - \tanh^{-1} \frac{u}{c}.$$

9. If  $M = \frac{mv}{\sqrt{1-v^2/c^2}}$ ,  $E = \frac{mc^2}{\sqrt{1-v^2/c^2}}$ ,  $F = \frac{dM}{dt}$ , prove that  $Fv = \frac{dE}{dt}$ ;  $m$  and  $c$  being constants, and  $v$  a function of  $t$ . Interpret these relations in connexion with special relativity.

## CHAPTER XIX

1. What is the significance of the so-called absolute determinations of the velocity of rotation of the earth by means of Foucault's pendulum or similar devices, in the general theory of relativity?

2. If we regard the earth as at rest, which according to the principle of relativity is allowable, how can we explain the rotation round the earth of distant stars in circular orbits with velocities enormously greater than that of light?

3. If in rectangular co-ordinates  $x, y, z, t$

$$ds^2 = -dx^2 - dy^2 - dz^2 + dt^2,$$

show that in a system of rectangular axes  $x_1, x_2, x_3, x_4$ , rotating with angular velocity  $\omega$  in the  $x, y$  plane,

$$ds^2 = -dx_1^2 - dx_2^2 - dx_3^2 + [1 - \omega^2(x_1^2 + x_2^2)]dx_4^2 + 2\omega x_2 dx_1 dx_4 - 2\omega x_1 dx_2 dx_4.$$

4. Show directly that the equations

$$\frac{\partial^2 x_\sigma}{\partial s^2} + \left\{ \alpha_\beta, \sigma \right\} \frac{\partial x_\alpha}{\partial s} \frac{\partial x_\beta}{\partial s} = 0 \quad (\sigma = 1, 2, 3, 4)$$

must be the equations of motion of a particle in a gravitational field.

5. Show that Einstein's law of gravitation may be regarded as an attraction inversely as the square of the distance together with a very small attraction inversely as the fourth power of the distance.

6. Show that the deflection of a particle moving with the velocity of light past the sun on Newton's law of gravitation is one-half that indicated by Einstein's theory.

7. Show that the deflection of light by the sun, on Einstein's theory, is the same as if the space around the sun had a refractive index  $1 + 2m/r$ , where  $m$  is the mass of the sun and  $r$  the distance from its centre.



# APPENDIX

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TABLE OF NUMERICAL VALUES

Velocity of light,	$2.9986 \times 10^{10}$ cm./sec.
Gravitation constant,	$6.66 \times 10^{-8}$ cm. <sup>3</sup> /gm. sec. <sup>2</sup> .
Protonic charge,	$e = 4.774 \times 10^{-10}$ e.s.u.
Electronic ratio of charge to mass,	$\frac{e}{m_0} = 1.769 \times 10^7$ e.m.u. per gramme.
Faraday,	9650 e.m.u.
Planck's constant,	$h = 6.554 \times 10^{-27}$ erg sec.
Gas constant,	$R = 8.315 \times 10^7$ erg/1° C. mol = 1.9869 calories/1° C.
Molecules per mol,	$6.061 \times 10^{23}$ .
Molecules per cubic centimetre in gas at 0° C. and 760 mm.,	$2.705 \times 10^{19}$ .
Gas constant for one molecule,	$k = 1.372 \times 10^{-16}$ erg/1° C.
Mass of hydrogen atom,	$1.663 \times 10^{-24}$ gm.
Mass of electron,	$m_0 = 8.999 \times 10^{-28}$ gm.
One atmosphere,	= 1,013,250 dynes per square centimetre.
Atomic weight of hydrogen,	1.0077.



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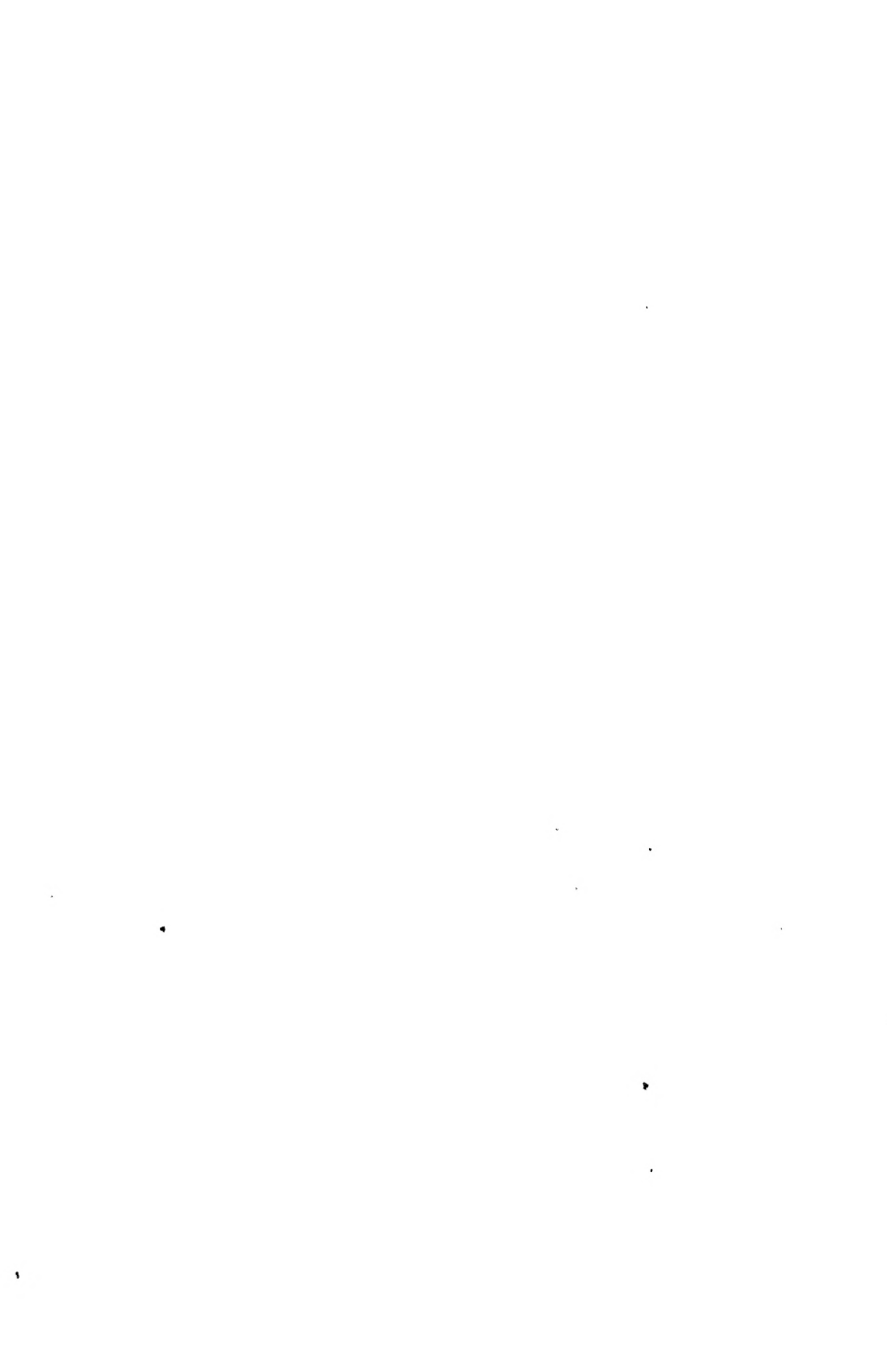
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